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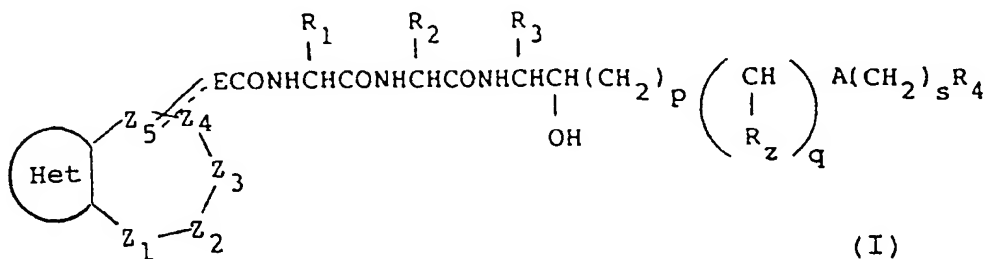
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(54) Novel compounds with renin-inhibiting activity.

(57) Compounds of formula (I), and pharmaceutically acceptable salts thereof:

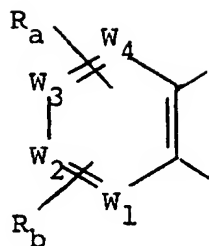


wherein
the Het ring is of sub-formula (a) or (b):

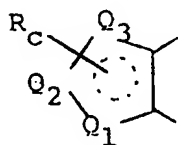
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USSN 10/619,769

PC25209A



(a)



(b)

wherein

one or two of W_1 , W_2 , W_3 and W_4 is N or NO (such that if two are NO then these are W_1 and W_4), and the others are CH, CR_a or CR_b ;

one of Q_1 , Q_2 and Q_3 is S and the other two are CH and CR_c ;

either Z_1 and Z_2 are absent and Z_3 , Z_4 and Z_5 and the carbon atoms to which Z_3 and Z_5 are attached, form a 5-membered non-aromatic heterocyclic ring; or

Z_1 is absent and Z_2 , Z_3 , Z_4 , Z_5 and the carbon atoms to which Z_2 and Z_5 are attached, form a 6-membered non-aromatic heterocyclic ring; or

Z_1 , Z_2 , Z_3 , Z_4 and Z_5 and the carbon atoms to which Z_1 and Z_5 are attached, form a 7-membered non-aromatic heterocyclic ring;

E is absent or is $(CH_2)_n$ or $CH(CH_2)_{n-1}$ wherein n is 1 to 4;

A is $-CONH-$, $-NHCO-$, $-COO-$, $-S(O)_r-$ wherein r is 0, 1 or 2, or $-CH_2-$;

p is 0, 1 or 2;

s is 0, 1, 2, 3 or 4;

q is 0 or 1;

R_2 is hydrogen, C_{1-6} alkyl or, when A is $-CH_2-$, hydroxy;

R_a and R_b are independently selected from hydrogen or a substituent;

R_1 is CH_2R_9 wherein R_9 is optionally substituted aryl or heteroaryl;

R_2 is $CHR_{10}R_{11}$ wherein R_{10} is hydrogen or methyl and R_{11} is C_{1-6} alkyl, C_{3-8} cycloalkyl, optionally substituted aryl or heteroaryl, or R_{11} is amino, C_{2-7} alkanoylamino, 2-oxopyrrolidinyl, 2-oxopiperidinyl or C_{1-6} alkoxy carbonylamino;

R_3 is CH_2R_{12} wherein R_{12} is C_{1-6} alkyl, C_{3-8} cycloalkyl or phenyl;

R_4 is C_{1-5} alkyl, C_{3-8} cycloalkyl, a saturated or unsaturated heterocyclic ring linked through carbon, hydroxy, C_{1-5} alkoxy, C_{1-7} alkanoyloxy, amino, C_{1-7} alkanoylamino, amino substituted by one or two C_{1-6} alkyl groups, or C_{1-6} alkylsulphonyl, carboxy, C_{1-6} alkoxy carbonyl, benzyloxycarbonyl, aminocarbonyl or $CH(NHR_{13})CO_2R_{14}$ wherein R_{13} is hydrogen or C_{1-6} alkanoyl and R_{14} is hydrogen or C_{1-6} alkyl; or (when s is 2 to 4) R_4 is a saturated or unsaturated heterocyclic ring linked through nitrogen; and

the dashed line represents an optional bond (when E is present);

which are renin inhibitors, a process for their preparation and their use as pharmaceuticals.

NOVEL COMPOUNDS

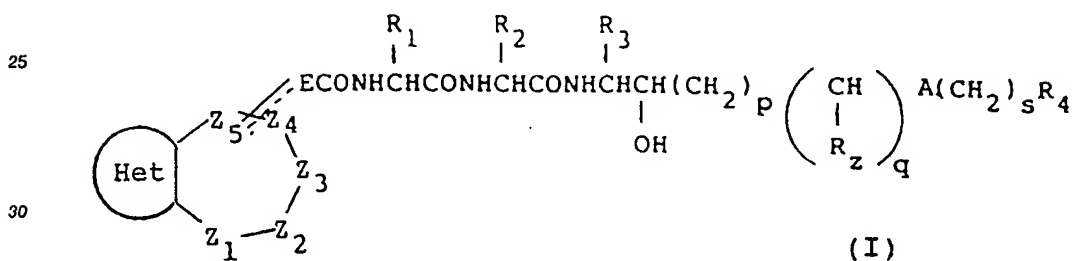
The invention relates to novel compounds having pharmacological activity, to processes for their preparation, to pharmaceutical preparations containing them and to their use in medicine.

Renin is a natural enzyme, disorders in relation to which are implicated in many cases of hypertension. It is released into the blood from the kidney, and cleaves from a blood glycoprotein a decapeptide known as angiotensin-I. Circulating angiotensin-I is cleaved in plasma, and in lung, kidney and other tissues to an octapeptide, angiotensin-II, which raises blood pressure both directly by causing arteriolar constriction and indirectly by stimulating release of the sodium-retaining hormone aldosterone from the adrenal gland and thus causing a rise in extracellular fluid volume. The latter effect is caused by angiotensin-II itself or a heptapeptide cleavage product angiotensin-III.

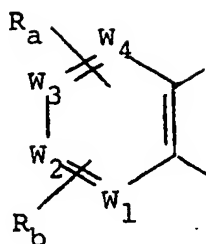
Inhibitors of renin have been described as useful in the treatment of hypertension.

A group of compounds has now been discovered, which inhibit the enzyme renin and therefore have potential blood pressure lowering activity, useful in the treatment of hypertension. Compounds having an associated mechanism of action have also been described as possessing anti-retroviral activity and the present compounds may therefore be of potential use in the treatment of diseases caused by retroviruses including human immunodeficiency virus (HIV-1 and 2) and HTLV I and II. They may also be of potential use in the treatment of other cardiovascular disorders, such as congestive heart failure, and have beneficial effects on learning and memory and mood elevation activity, of potential use in the treatment of CNS disorders such as Alzheimer's disease and depression; they may also be of potential use in the treatment of glaucoma.

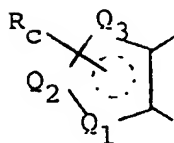
Accordingly, the present invention provides a compound of formula (I), or a pharmaceutically acceptable salt thereof:



wherein
the Het ring is of sub-formula (a) or (b):



(a)



(b)

wherein

one or two of W_1 , W_2 , W_3 and W_4 is N or NO (such that if two are NO then these are W_1 and W_4), and the others are CH, CR_a or CR_b ;

one of Q_1 , Q_2 and Q_3 is S and the other two are CH and CR_c ;

either Z_1 and Z_2 are absent and Z_3 , Z_4 and Z_5 and the carbon atoms to which Z_3 and Z_5 are attached, form a 5-membered non-aromatic heterocyclic ring; or

Z_1 is absent and Z_2 , Z_3 , Z_4 , Z_5 and the carbon atoms to which Z_2 and Z_5 are attached, form a 6-membered non-aromatic heterocyclic ring; or

Z_1 , Z_2 , Z_3 , Z_4 and Z_5 and the carbon atoms to which Z_1 and Z_5 are attached, form a 7-membered non-aromatic heterocyclic ring;

E is absent or is $(CH_2)_n$ or $CH(CH_2)_{n-1}$ wherein n is 1 to 4;

A is -CONH-, -NHCO-, -COO-, -S(O)r- wherein r is 0, 1 or 2, or -CH₂-;

p is 0, 1 or 2;

s is 0, 1, 2, 3 or 4;

q is 0 or 1;

R_2 is hydrogen, C₁₋₆ alkyl or, when A is -CH₂-, hydroxy;

R_a and R_b are independently selected from hydrogen or a substituent;

R_1 is CH_2R_9 wherein R_9 is optionally substituted aryl or heteroaryl;

R_2 is $CHR_{10}R_{11}$ wherein R_{10} is hydrogen or methyl and R_{11} is C₁₋₆ alkyl, C₃₋₈ cycloalkyl, optionally substituted aryl or heteroaryl, or R_{11} is amino, C₂₋₇ alkanoylamino, 2-oxopyrrolidinyl, 2-oxopiperidinyl or C₁₋₅ alkoxy-carbonylamino;

R_3 is CH_2R_{12} wherein R_{12} is C₁₋₆ alkyl, C₃₋₈ cycloalkyl or phenyl;

R_4 is C₁₋₆ alkyl, C₃₋₈ cycloalkyl, a saturated or unsaturated heterocyclic ring linked through carbon, hydroxy, C₁₋₅ alkoxy, C₁₋₇ alkanoyloxy, amino, C₁₋₇ alkanoylamino, amino substituted by one or two C₁₋₆ alkyl groups, or C₁₋₆ alkylsulfonyl, carboxy, C₁₋₆ alkoxy-carbonyl, benzyloxy-carbonyl, aminocarbonyl or $CH(NHR_{13})CO_2R_{14}$ wherein R_{13} is hydrogen or C₁₋₆ alkanoyl and R_{14} is hydrogen or C₁₋₆ alkyl; or (when s is 2 to 4) R_4 is a saturated or unsaturated heterocyclic ring linked through nitrogen; and the dashed line represents an optional bond (when E is present).

When the Het ring is of sub-formula (a)

Often W_3 or W_4 is N or NO, usually W_3 is N when E is attached at Z_5 , or W_1 is N when E is attached at Z_5 .

5 When the Het ring is of sub-formula (b)

Often Q_1 or Q_3 is S.

10 When Z_1 and Z_2 are absent:

Values of Z_3 , Z_4 and Z_5 include wherein one of Z_3 , Z_4 and Z_5 is attached to E and is N, CH or C (wherein the optional exocyclic bond is present); the second of Z_3 , Z_4 and Z_5 is O, S, SO, SO_2 or NR wherein R is hydrogen or C_{1-6} alkyl; and the third is O, S, SO, SO_2 , NR, CH_2 or C=O provided that, one of
 15 Z_3 and Z_4 , and one of Z_4 and Z_5 , is CH_2 , CH or C when the other is S or O, or is CH_2 , CH, C, NR or N when the other is SO or SO_2 ; or Z_4 is SO, SO_2 or C=O when one of Z_3 and Z_5 is N and the other is O, S or NR; or Z_3 is CO, Z_4 is O and Z_5 is CH.

Examples of Z_3 , Z_4 and Z_5 then include

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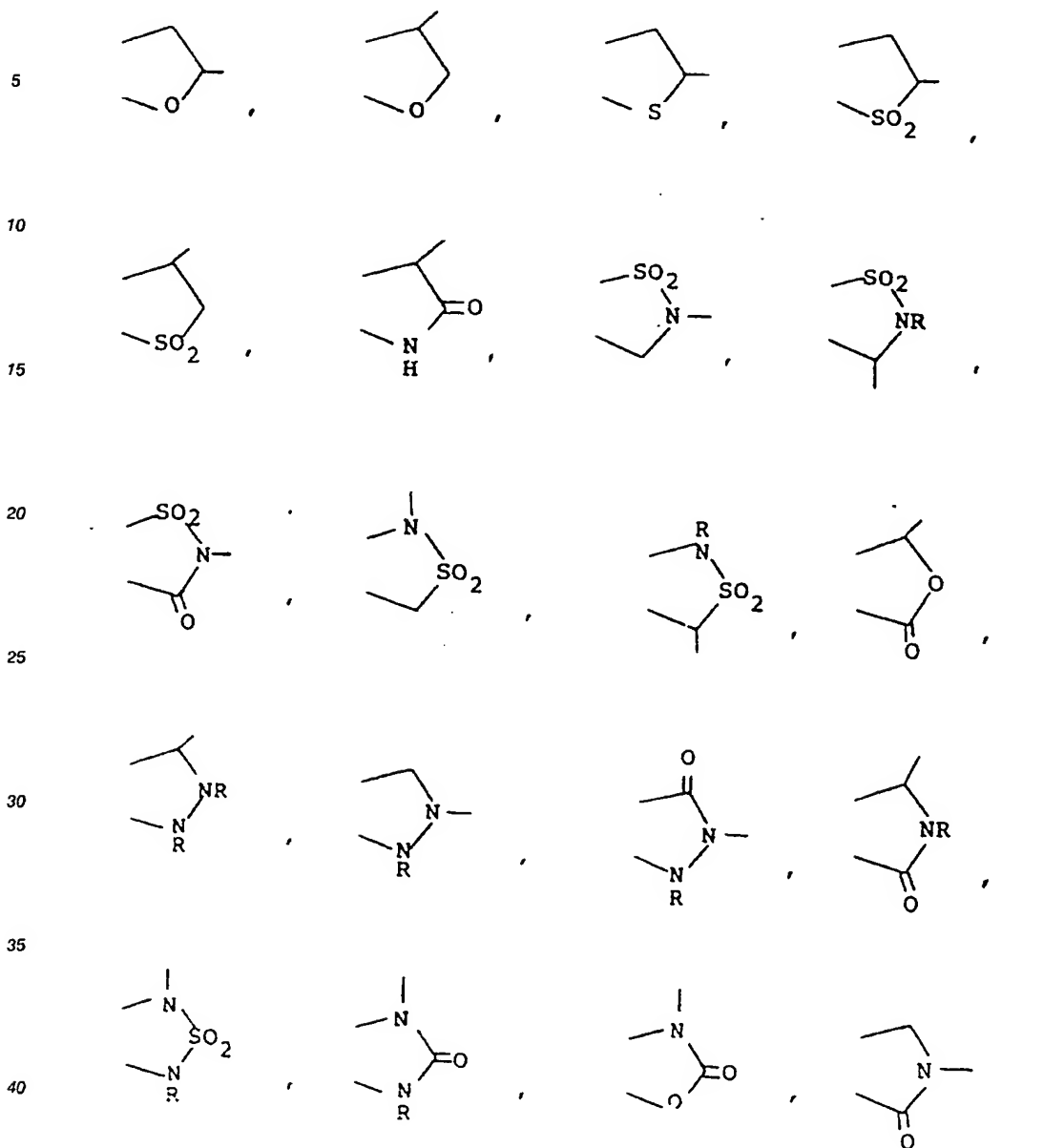
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45 Preferably Z_3 is CO, Z_4 is N, Z_5 is CH and E is attached at Z_4 .

When Z_1 is absent:

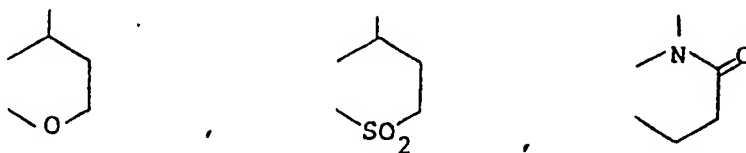
50 Values of Z_2 , Z_3 , Z_4 and Z_5 include wherein one of Z_2 , Z_3 , Z_4 and Z_5 is attached to E and is N, CH or C (wherein the optional exocyclic bond is present); another of Z_2 , Z_3 , Z_4 and Z_5 is O, S, SO, SO₂ or NR wherein R is hydrogen or C₁₋₆ alkyl; another of Z_2 , Z_3 , Z_4 and Z_5 is O, S, SO, SO₂, NR or CH₂; and the other of Z_2 , Z_3 , Z_4 and Z_5 is S, SO, SO₂, NR, CH₂ or C=O provided that, one of Z_2 and Z_3 , one of Z_3 and Z_4 , or one of Z_4 and Z_5 is CH₂, CH or C when the other is S or O, or is CH₂, CH, C, NR or N when the other is SO or SO₂; or Z_3/Z_4 is SO, SO₂ or C=O when one of Z_2/Z_3 and Z_4/Z_5 is N or NR and the other is O, S, N or NR; or $Z_2-Z_3-Z_4-Z_5$ is -NRCOCONR- or >NCOCONR-; or Z_2/Z_5 is CO, Z_3/Z_4 is O and $Z_4, Z_5/Z_2, Z_3$ are CH and CH₂.

i.e. the following are excluded in Z_2 to Z_5 :-

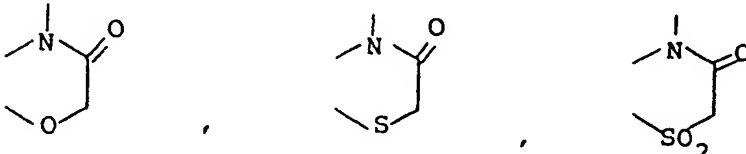
- i) adjacent same heteroatoms (O, S, N) except N-N; but no N-N-N;
- iii) $\text{SO}_x\text{-SO}_y$ or O-S_y (x and y are 0-2);
- iv) O-C-O , S-C-O , S-C-S ;
- v) CO-SO or CO-SO_2 ;
- 5 vi) two heteroatoms (O, S, N) separated by a Z when CO unless one of them is N;
- vii) more than two of Z_2 to Z_5 is CO.

Examples of Z_2 , Z_3 , Z_4 and Z_5 then include

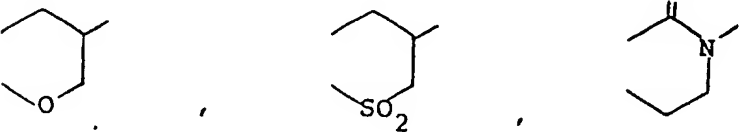
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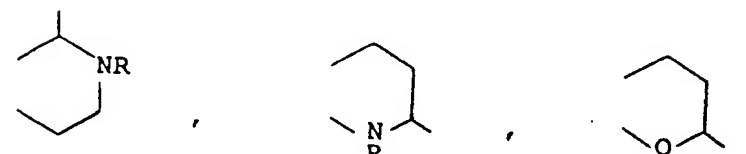


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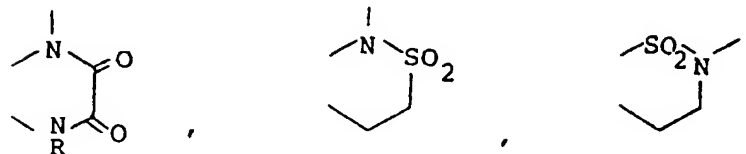


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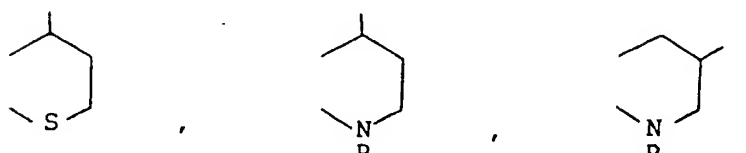
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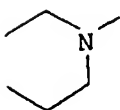


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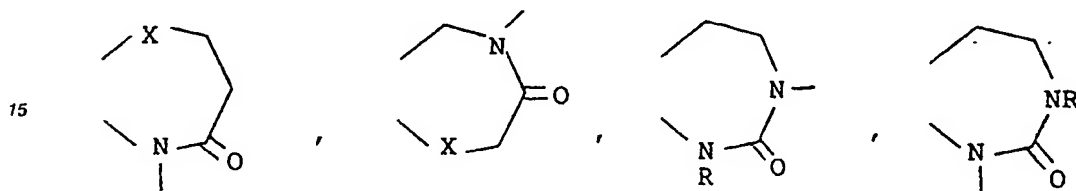
When W_3 is N and the other W are CH, CR_a or CR_b , often Z_2 is S, Z_3 is CH_2 , Z_4 is CO, Z_5 is N and E is attached at Z_5 .

5 When Z_1 and Z_2 are present

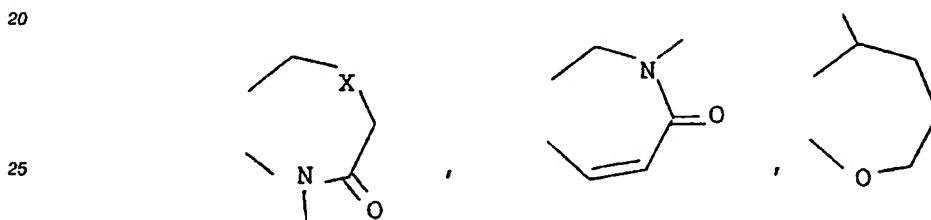
Heterocyclic ring heteroatoms for Z_1 to Z_5 are selected from oxygen, sulphur, SO, SO_2 , nitrogen or N-alkyl. The ring may include unsaturation ($C=C$) or a carbon attached to exocyclic oxo ($C=O$).

Examples of Z_1 , Z_2 , Z_3 , Z_4 and Z_5 then include

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30 wherein X is O, S, SO, SO_2 , NH, $N-C_{1-6}$ alkyl or CH_2 .

Preferably E is attached at a nitrogen atom.

Suitable examples of R_a and R_b in sub-formula (a), include a moiety selected from halogen, C_{1-6} alkyl, C_{1-6} alkoxy, cyano, CHO, C_{1-6} alkyl substituted by optionally substituted amino or R_a/R_b is a group NR_5R_6 , CO_2R_5 , wherein R_5 is hydrogen, C_{1-6} alkyl or optionally substituted benzyl and R_6 is hydrogen or C_{1-6} alkyl.

When one of R_a and R_b is hydrogen and the other is a halogen substituent, the substituent is preferably in the 3-position relative to the ring nitrogen atom, when one of W_1 to W_4 is N or NO.

Suitable examples of R_c in sub-formula (b), include a moiety selected from C_{1-6} alkyl, optionally substituted amino, cyano, C_{1-6} alkyl substituted by optionally substituted amino or R_c is CO_2R_5 wherein R_5 is hydrogen, C_{1-6} alkyl or optionally substituted benzyl.

Suitable examples of substituents in optionally substituted amino groups in $R_a/R_b/R_c$ then include one or two groups independently selected from C_{1-6} alkyl or C_{1-6} alkyl substituted by carboxy or C_{1-6} alkoxy carbonyl.

Alternatively, NR_5R_6 or other substituted amino groups in $R_a/R_b/R_c$ may be pyrrolidinyl, piperidinyl, morpholinyl or piperazinyl; optionally N-substituted by C_{1-6} alkyl, or as an N-oxide thereof.

Preferably one of R_a and R_b is hydrogen and the other is CH_2NH_2 , CO_2H or $CH_2NHCH_2CO_2H$.

R_c is preferably hydrogen.

In regard to values of A and p and q, reference is hereby made to the following literature:-

1. J. Med. Chem. 1988, 31, 1918
 (p = 1, q = 0, A = $-CONH-$)
 2. J. Med. Chem. 1987, 30, 1224
 (p = 2, q = 0, A = $-NHCO-$)
 3. J. Med. Chem. 1987, 31, 701
 (p = 0, q = 0, A = $-COO-$)
 4. J. Med. Chem. 1987, 31, 1839
 (p = 1, q = 1, A = $-CONH-$)
 5. J. Med. Chem. 1987, 30, 1729 and refs. therein
 (p = 2, q = 1, A = $S(O)_n$)

6. WO 88/05050

(p = 0, q = 1, A = CH₂, R_z = OH)

7. EP-A-172346

(p = 0, q = 1, A = CH₂, R_z = H)

- 5 Suitable values of R₉ and R₁₁ when aryl include phenyl or naphthyl and when heteroaryl, include a 5- or 6-membered monocyclic or 9- or 10- membered bicyclic of which 5- or 6- membered monocyclic heteroaryl is preferred. In addition, 5- or 6-membered monocyclic or 9- or 10-membered bicyclic heteroaryl preferably contains one, two or three heteroatoms which are selected from the class of oxygen, nitrogen and sulphur and which, in the case of there being more than one heteroatom, are the same or different.
- 10 Examples of 5- or 6-membered monocyclic heteroaryl containing one, two or three heteroatoms which are selected from the class of oxygen, nitrogen and sulphur include furyl, thienyl, pyrrolyl, oxazolyl, thiazolyl, imidazolyl and thiadiazolyl, and pyridyl, pyridazyl, pyrimidyl, pyrazolyl and triazolyl. Preferred examples of such groups include furanyl, thienyl, pyrrolyl and pyridyl, in particular 2- and 3-furanyl, 2- and 3-pyrrolyl, 2- and 3-thienyl, and 2-, 3- and 4-pyridyl. Examples of 9- or 10-membered bicyclic heteroaryl containing one, two
- 15 or three heteroatoms which are selected from the class of oxygen, nitrogen and sulphur include benzofuranyl, benzothienyl, indolyl and indazolyl, quinolyl and isoquinolyl, and quinazolyl. Examples of such groups include 2- and 3-benzofuranyl, 2- and 3-benzothienyl, and 2- and 3-indolyl, and 2- and 3-quinolyl and, for R₁₁, 4-benzimidazolyl.

- Suitable examples of groups or atoms for optional substitution of R₅ when benzyl and R₉ and R₁₁ include one, two or three substituents independently selected from C₁₋₄ alkyl, C₁₋₄ alkoxy, halo (such as fluoro, chloro, bromo), hydroxy, nitro, and cyano.
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Preferred examples of R₉ include phenyl and naphthyl, and preferred examples of R₁₁ when aryl or heteroaryl include phenyl, imidazol-4-yl and -2-yl, pyrazol-1-yl and 4-methylpyrazol-1-yl.

Preferably the amino acid residue containing R₂ is Leu, β-pyrazolylalanine or His.

- 25 Abbreviations which may be used herein are as follows:

Amino Acid Residue	Abbreviations
histidine	His
isoleucine	Ile
leucine	Leu
1-naphthylalanine	NAla
norleucine	Nle
phenylalanine	Phe

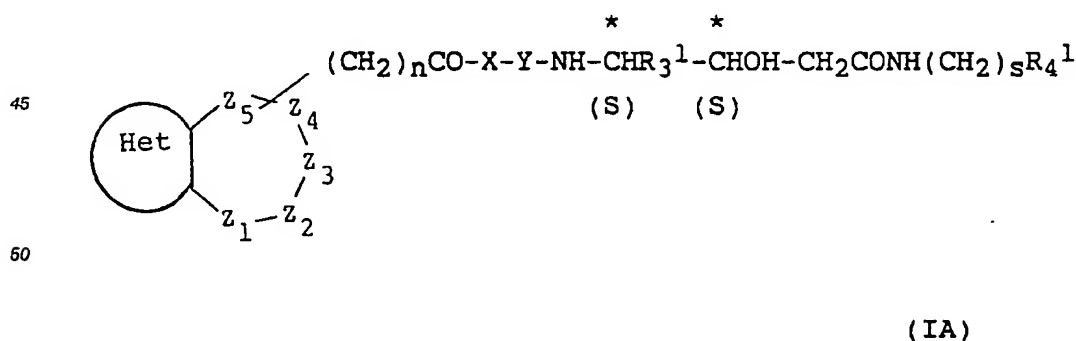
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The α-amino acid components are in the (S)-configuration (or L-form).

In a particular aspect, the present invention provides a compound of formula (IA) or a pharmaceutically acceptable salt thereof:

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wherein

X is Phe;

Y is Leu or His;

R₃' is cyclohexylmethyl;

R₄' is C₄₋₅ alkyl and s is 0 or s is 2 to 4 and R₄' is carboxy or a saturated or unsaturated heterocyclic ring; and

the remaining variables are as defined in formula (I).

- 5 Suitable values for alkyl groups in R_a, R_b, R₂, R and R₂ to R₁₂ in formulae (I) and (IA) include methyl, ethyl, n - and iso -propyl, n -, sec -, iso - and tert -butyl, n -, iso -, sec -, tert - and neo -pentyl. C₃₋₈ cycloalkyl groups are cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl and cyclooctyl.

Suitable values for R_a and R_b halo include fluoro, chloro, bromo and iodo, preferably chloro or bromo.

Preferably R₄ or R₄' when alkyl is iso -butyl.

- 10 Suitable examples of R₄ when monocyclic heteroaryl include those listed for R₉ and R₁₁, preferably 3-pyridyl or an N-oxide thereof, or 1- or 2-imidazolyl.

When R₄ is a saturated heterocyclic ring, suitable examples include piperidinyl, pyrrolidinyl, piperazinyl and morpholinyl; optionally N-substituted by C₁₋₅ alkyl group(s) or as an N-oxide thereof.

- 15 The right hand sequence in formula (I) and (IA) is preferably 4(S)-amino-5-cyclohexyl-3(S)-hydroxyphen-tanoic acid (ACHPA), from the amino acid containing R₃ or R₃' to NH(CH₂)₅R₄.

Preferably n is 1, 2, 3 or 4.

- When Z₁ is absent, E is favourably attached at the 4-position with respect to Z₂ when n=2, and at the 3-position when n>2. When Z₁ and Z₂ are absent, E is preferably at the 3-position with respect to Z₃ when n=2, and at the 2-position when n>2. When Z₁ and Z₂ are present, E is usually attached at a nitrogen atom
20 or at Z₁ or Z₅ when a carbon atom. E is preferably attached at a nitrogen atom.

- Pharmaceutically acceptable salts include acid addition salts which may be, for example, salts with inorganic acids such, for example, as hydrochloric acid, hydrobromic acid, orthophosphoric acid or sulphuric acid, or organic acids such, for example, as methanesulphonic acid, toluenesulphonic acid, acetic acid, trifluoroacetic acid, propionic acid, lactic acid, citric acid, tartaric acid, fumaric acid, malic acid,
25 succinic acid, salicylic acid or acetylsalicylic acid.

Pharmaceutically acceptable salts may also include alkali metal salts, for example sodium or potassium, alkaline earth metal salts such as calcium or magnesium and ammonium or substituted ammonium salts, for example those with lower alkylamines such as triethylamine, hydroxy-lower alkylamines such as 2-hydroxyethylamine, bis-(2-hydroxyethyl)-amine or tris-(2-hydroxyethyl)-amine.

- 30 It will be appreciated that the compounds of the invention may exist as solvates, such as hydrates, and these are included whenever, a compound of formula (I) or a salt thereof, is herein referred to.

- The compounds of formula (I) wherein E is attached to Z₁(when present)/Z₂(when present)/Z₃/Z₄/Z₅ when CH, and (IA) have at least one asymmetric centre in addition to those attached to R₁ and R₂ (in X and Y) and R₃ and those indicated in formula (IA), and are therefore capable of existing in more than one
35 stereoisomeric form. The invention extends to each of these forms and to mixtures thereof. Preferably the configuration at the carbon atoms bearing R₁, R₂ and R₃ is the (S)-configuration.

It will be appreciated that, when the optional bond depicted in formula (I) is present, the compounds are capable of existing in E and Z (trans and cis) forms.

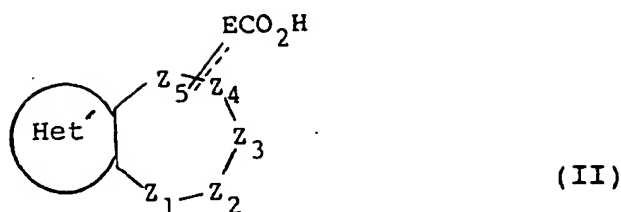
The invention extends to each of these forms and to mixtures thereof.

- 40 The compounds of the invention are preferably in pharmaceutically acceptable form. By pharmaceutically acceptable form is meant, inter alia, of a pharmaceutically acceptable level of purity excluding normal pharmaceutical additives such as diluents and carriers, and including no material considered toxic at normal dosage levels. A pharmaceutically acceptable level of purity will generally be at least 50% excluding normal pharmaceutical additives, preferably 75%, more preferably 90% and still more preferably 95%.

- 45 A compound of the invention may be prepared by those methods known in the art for the synthesis of compounds of analogous structure, such as peptides, and in this regard reference is made, by way of illustration only, to the literature reference: S.R. Pettit, " Synthetic Peptides ", (Elsevier Scientific Publishing Co. 1976).

- The present invention also provides a compound of the present invention which has been prepared
50 synthetically.

A compound of the present invention may, for example, be formed by the sequential coupling of appropriate amino acids with the acid of formula (II):

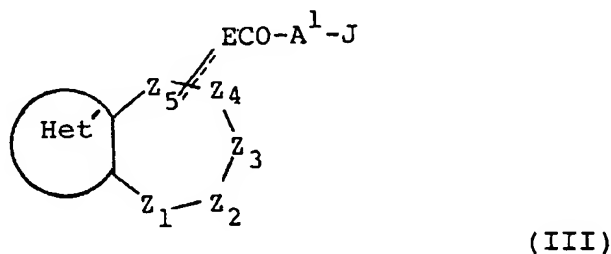


10 wherein Het' is Het as defined in formula (I), but substituted by R_a' , R_b' in sub-formula (a) and by R_c' in sub-formula (b), wherein R_a' , R_b' and R_c' are R_a , R_b and R_c respectively or groups or atoms convertible thereto; or by the initial preparation and subsequent coupling of peptide subunits with the acid of formula (II), the subunits themselves prepared in stepwise manner; in either case classical solution chemistry methods analogous to those used in peptide synthesis, may be employed.

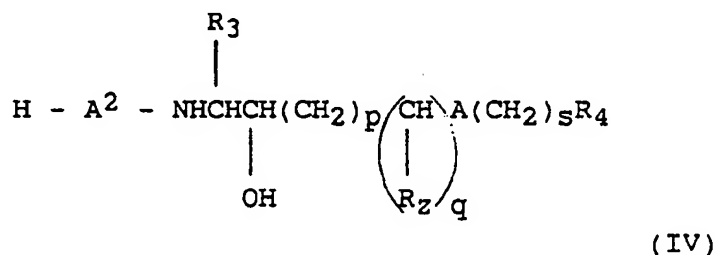
15 The coupling reactions may be effected by, for example, activating the reacting carboxyl group of the acid of formula (II) or amino acid, and reacting this with the amino group of the substrate unit. Details of suitable, optional activating and protecting (masking) groups and of suitable reaction conditions (for the coupling reactions and for the introduction and removal of protecting groups) giving, preferably, the minimum of racemisation, may be found in the above-referenced literature.

20 It will be appreciated that, when A is other than -CONH- or -COO-, the C-terminus group is other than an amino acid, in which case the coupling is carried out with an appropriate other precursor, as described in the aforementioned literature -references 2., 5., 6. and 7.

25 Accordingly, the present invention further provides a process for the preparation of a compound of formula (I) or a pharmaceutically acceptable salt thereof which comprises reacting a reagent of the formula (III):



35 wherein Het' is Het as defined but wherein R_a , R_b and R_c are R_a' , R_b' and R_c' , which are R_a , R_b and R_c respectively or groups or atoms convertible thereto; A^1 is absent or represents an appropriate amino acid or dipeptide unit; J is OH or a leaving group; and the remaining variables are as hereinbefore defined; with a reagent of the formula (IV):



55

wherein

A^2 is absent or represents an appropriate amino acid or dipeptide unit, such that $A^1 + A^2$ is -NHCHR₁CONHCHR₂CO-, and the remaining variables are as hereinbefore defined; and thereafter, if

desired or necessary deprotecting (within A¹ or A² of the products, and/or converting Z₁, Z₂, Z₃, Z₄ or Z₅ to other Z₁, Z₂, Z₃, Z₄ or Z₅, R_a/R_b/R_c in Het' to R_a, R_b and/or R_c and/or W₁, W₂, W₃ or W₄ in Het' when N to NO; and/or forming a pharmaceutically acceptable salt thereof.

Suitable examples of J when a leaving group include halo, such as chloro or bromo, and other suitable groups which are displaceable by an amino nucleophile, such as C₁₋₆ alkoxycarbonyloxy.

It is generally preferred, especially when A¹ is not absent, however, that J is OH, and a suitable coupling reagent or dehydrating catalyst is used to effect the reaction, such as N,N-dicyclohexylcarbodiimide and those described in the Descriptions and Examples hereinafter.

Deprotection of A¹ and/or A² takes place conventionally, in accordance with the particular protecting group(s) employed.

Pharmaceutically acceptable salts may be formed conventionally.

Z₁, Z₂, Z₃, Z₄ or Z₅ when S or SO may be converted to SO or SO₂ respectively (or S to SO₂) by conventional oxidation methods, preferably in a non-aqueous solvent, such as a chlorinated hydrocarbon, in the presence of an organic peracid, such as 3-chloroperoxybenzoic acid, or in water in the presence of a soluble strong inorganic oxidant, such as an alkali metal permanganate or with aqueous hydrogen peroxide. A particularly suitable alternative oxidation method involves the use of magnesium monopero-phthalate, as described in Description 10b) hereinafter.

It will be apparent that compounds of the formula (I) containing a Het' ring wherein the R_a'/R_b'/R_c' group is other than R_a/R_b/R_c and which is convertible to an R_a/R_b/R_c group, are useful novel intermediates. A number of such conversions is possible, not only for the final product compounds of formula (I) when R_a'/R_b'/R_c' are other than R_a/R_b/R_c, but also within R_a/R_b/R_c, and also for their intermediates as follows:

- (i) an R_a/R_b nitro substituent is convertible to an amino substituent by reduction;
- (ii) a C₁₋₇ acylamino substituent is convertible to an amino substituent by deacylation;
- (iii) a hydrogen substituent in sub-formula (a) is convertible to a halogen substituent by halogenation;
- (iv) an amino substituent is convertible to a corresponding substituent which is substituted by one or two alkyl groups, by N-alkylation;
- (v) a cyano substituent may be converted to an aminomethyl substituent by reduction;
- (vi) a bromo substituent may be converted to a cyano substituent by reaction with copper(I) cyanide.

Conversions (i) to (vi) are only exemplary and are not exhaustive of the possibilities. It will be appreciated that it is often desirable to carry out these conversions at an earlier stage, in the intermediate acid of formula (II).

In regard to (i), the reduction is carried out with a reagent suitable for reducing nitroanisole to aminocanisole.

In regard to (ii), deacylation is carried out by treatment with a base, such as an alkali metal hydroxide.

In regard to (iv), alkylation is carried out with a corresponding alkylating agent such as the chloride or bromide under conventional conditions.

In regard to (v), the reduction takes place by reaction with sodium borohydride/cobalt chloride or platinum oxide/hydrogen.

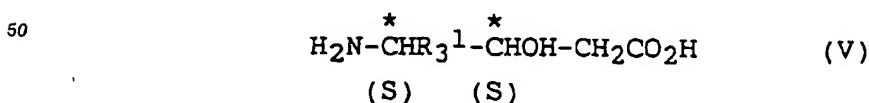
In regard to (vi), the reaction takes place under conventional conditions.

Conversion of W₁, W₂, W₃ or W₄ when N to NO may be achieved by similar oxidation methods to those used for Z₁, Z₂, Z₃, Z₄ or Z₅ when S or SO to SO or SO₂, such as using 3-chloroperoxybenzoic acid. Selective oxidation may be achieved by an appropriate choice of oxidising agent and conditions.

Compounds of the general formulae (II) and (III) may themselves be prepared by standard techniques analogous to those described above.

The acids of formula (II) are either known compounds or are prepared by analogous methods to these and for structurally similar known compounds.

The preparation of the amino acid of formula (V):



is described in J. Med. Chem 1985, 28, 1779-1790.

As regards the appropriate corresponding intermediates to those of formula (V), including those wherein A is other than -CONH-, reference is hereby made to the literature references 2. to 7. listed hereinbefore.

It will be appreciated that protected forms of compounds of the present invention are novel inter-

mediates and form an aspect of the invention.

Particularly suitable methods of preparation of the compounds of the present invention are as described in the Descriptions and Examples hereinafter. The couplings may be carried out sequentially beginning by coupling the acid of formula (II) with, for example, phenylalanine or 1-naphthylalanine, followed by coupling
 5 with Y, for example, leucine or histidine, and finally, coupling with the amino acid of formula (V). In a preferred aspect, however, either the acid of formula (II) is coupled with a tripeptide unit formed between the amino acid of formula (V); leucine, histidine or other R₂ containing amino acid; and phenylalanine or naphthylalanine; or alternatively the acid of formula (II) is coupled with phenylalanine or naphthylalanine and this is coupled with a dipeptide unit formed between the R₂ containing amino acid and the amino acid of
 10 formula (V).

As mentioned previously the compounds of the invention have been found to be renin inhibitors, and therefore they are of potential use in the treatment of hypertension. They may also be of potential use in the other diseases and disorders hereinbefore referred to.

The present invention also provides a pharmaceutical composition which comprises a compound of
 15 formula (I) or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier. In particular, the present invention provides an anti-hypertensive pharmaceutical composition which comprises an anti-hypertensive effective amount of a compound of formula (I) or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.

The compositions are preferably adapted for oral administration. However, they may be adapted for
 20 other modes of administration, for example parenteral administration for patients suffering from heart failure. Other alternative modes of administration include sublingual or transdermal administration, or for example, eye-drop compositions, for the treatment of glaucoma.

The compositions may be in the form of tablets, capsules, powders, granules, lozenges, suppositories, reconstitutable powders, or liquid preparations, such as oral or sterile parenteral solutions or suspensions.

25 In order to obtain consistency of administration it is preferred that a composition of the invention is in the form of a unit dose.

Unit dose presentation forms for oral administration may be tablets and capsules and may contain conventional excipients such as binding agents, for example syrup, acacia, gelatin, sorbitol, tragacanth, or polyvinylpyrrolidone; fillers, for example lactose, sugar, maize-starch, calcium phosphate, sorbitol or glycine;
 30 tableting lubricants, for example magnesium stearate; disintegrants, for example starch, polyvinylpyrrolidone, sodium starch glycolate or microcrystalline cellulose; or pharmaceutically acceptable wetting agents such as sodium lauryl sulphate.

The solid oral compositions may be prepared by conventional methods of blending, filling or tableting. Repeated blending operations may be used to distribute the active agent throughout those compositions
 35 employing large quantities of fillers. Such operations are of course conventional in the art. The tablets may be coated according to methods well known in normal pharmaceutical practice, in particular with an enteric coating.

Oral liquid preparations may be in the form of, for example, emulsions, syrups, or elixirs, or may be presented as a dry product for reconstitution with water or other suitable vehicle before use. Such liquid
 40 preparations may contain conventional additives such as suspending agents, for example sorbitol, syrup, methyl cellulose, gelatin, hydroxyethylcellulose, carboxymethylcellulose, aluminium stearate gel, hydrogenated edible fats; emulsifying agents, for example lecithin, sorbitan monooleate, or acacia; non-aqueous vehicles (which may include edible oils), for example almond oil, fractionated coconut oil, oily esters such as esters of glycerine, propylene glycol, or ethyl alcohol; preservatives, for example methyl or propyl p-
 45 hydroxybenzoate or sorbic acid; and if desired conventional flavouring or colouring agents.

For parenteral administration, fluid unit dosage forms are prepared utilizing the compound and a sterile vehicle, and, depending on the concentration used, can be either suspended or dissolved in the vehicle. In preparing solutions the compound can be dissolved in water for injection and filter sterilized before filling
 50 into a suitable vial or ampoule and sealing. Advantageously, adjuvants such as a local anaesthetic, a preservative and buffering agents can be dissolved in the vehicle. To enhance the stability, the composition can be frozen after filling into the vial and the water removed under vacuum. Parenteral suspensions are prepared in substantially the same manner, except that the compound is suspended in the vehicle instead of being dissolved, and sterilization cannot be accomplished by filtration. The compound can be sterilized by exposure to ethylene oxide before suspending in the sterile vehicle. Advantageously, a surfactant or
 55 wetting agent is included in the composition to facilitate uniform distribution of the compound.

For transdermal administration, formulations suitable for topical use may be employed, optionally containing penetration enhancers.

The compositions may contain from 0.1% to 99% by weight, preferably from 10-60% by weight, of the

active material, depending on the method of administration.

The present invention further provides a method of prophylaxis or treatment of hypertension in mammals including man, which comprises administering to the suffering mammal an anti-hypertensively effective amount of a compound of formula (I) or a pharmaceutically acceptable salt thereof.

5 An effective amount will depend on the relative efficacy of the compounds of the present invention, the severity of the hypertension being treated and the weight of the sufferer. However, a unit dose form of a composition of the invention may contain from 0.1 to 500 mg of a compound of the invention and more usually from 1 to 100 mg, for example 2 to 50 mg such as 2, 3, 4, 5, 10, 20 or 30mg. Such compositions may be administered from 1 to 6 times a day, more usually from 2 to 4 times a day, in a manner such that
10 the daily dose is from 1 to 1000mg for a 70 kg human adult and more particularly from 5 to 500 mg.

No toxicological effects are indicated at the aforementioned dosage ranges.

The present invention further provides a compound of formula (I) or a pharmaceutically acceptable salt thereof for use in the treatment or prophylaxis of hypertension.

15 The following Descriptions relate to the preparation of intermediates and the following Examples relate to the preparation of compounds of formula (I).

The following tabulations show the structures prepared.

(* indicates the point of attachment of E).

In the following, the abbreviations used are:

20

Abbreviation	Definition
ACHPAA	4(S)-amino-5-cyclohexyl-3(S)-hydroxypentanoic acid isobutylamide
ACHPA	4(S)-amino-5-cyclohexyl-3(S)-hydroxypentanoic acid
BOC	tert-butoxycarbonyl
25 CBZ	benzyloxycarbonyl
ⁱ Bu	isobutyl
API	3-(1-imidazolyl)propylamino

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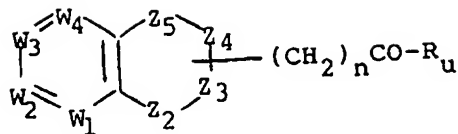
TABLE 1

35

R _s -ACHPA-R _t			
Desc.No.	R _s	R _t	salt/solvate
D3(a)	BOC	API	
D3(b)	CBZ-Phe-Leu	API	
D3(c)	Phe-Leu	API	
40 D4(a)	BOC-Phe-Leu	NH ⁱ Bu	CF ₃ CO ₂ H
D4(b)	Phe-Leu	NH ⁱ Bu	
D5(a)	CBZ-Phe-Leu	NH ⁱ Bu	
D5(b)	Phe-Leu	NH ⁱ Bu	CH ₃ CO ₂ H
D5(c)	Phe-Leu	NH ⁱ Bu	
45 D9(a)	BOC-Phe-Leu	API	
D9(b)	Phe-Leu	API	2CF ₃ CO ₂ H

50

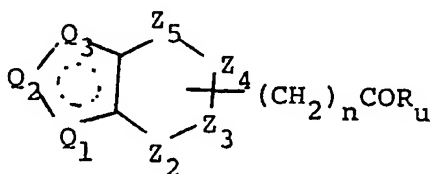
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TABLE 2

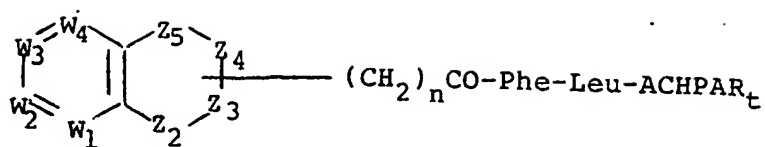
Desc.No.	W ₁	W ₂	W ₃	W ₄	Z ₂	Z ₃	Z ₄	Z ₅	n	R _u
D1	CH	CH	CH	N	-	O	CH ₂	CH*	1	OH
D2(a)	CH	CH	CH	N	O	CH ₂	CO	N*	2	OMe
D2(b)	CH	CH	CH	N	O	CH ₂	CO	N*	2	OH
D6(b)	CH	CH	N	CH	S	CH ₂	CO	N*	2	OMe
D6(b)	CH	CH	N	CH	S	CH ₂	CO	N*	2	OH
D7(c)	CH	CH	CH	N	-	NH	CO	N*	2	OEt
D7(d)	CH	CH	CH	N	-	NH	CO	N*	2	OH
D8(c)	N	CH	CH	CH	-	NH	CO	N*	2	OMe
D8(d)	N	CH	CH	CH	-	NH	CO	N*	2	OH
D10(a)	CH	CH	N	CH	S	CH ₂	CO	N*	2	O ^t Bu
D10(b)	CH	CH	N	CH	SO ₂	CH ₂	CO	N*	2	O ^t Bu
D10(c)	CH	CH	N	CH	SO ₂	CH ₂	CO	N*	2	OH
D11(b)	N	CH	CH	CH	S	CH ₂	CO	N*	2	O ^t Bu
D11(c)	N	CH	CH	CH	S	CH ₂	CO	N*	2	OH
D12(a)	N	CH	CH	CH	SO ₂	CH ₂	CO	N*	2	O ^t Bu
D12(b)	N	CH	CH	CH	SO ₂	CH ₂	CO	N*	2	OH
D13(a)	N	CH	CH	CH	-	CO	N*	CO	3	OEt
D13(b)	N	CH	CH	CH	-	CH(OH)	N*	CO	3	OEt
D13(c)	N	CH	CH	CH	-	CH ₂	N*	CO	3	OEt
D13(d)	N	CH	CH	CH	-	CH ₂	N*	CO	3	OH

TABLE 2 (Cont'd)

Desc.No.	W ₁	W ₂	W ₃	W ₄	Z ₂	Z ₃	Z ₄	Z ₅	n	R _u
D14(a)	N	CH	CH	CH	-	CO	N*	CH(OH)	3	OEt
D14(b)	N	CH	CH	CH	-	CO	N*	CH ₂	3	OEt
D14(c)	N	CH	CH	CH	-	CO	N*	CH ₂	3	OH
D15(a)	CH	N	CH	CH	-	CH	N*	CO	3	OEt
D15(b)	CH	N	CH	CH	-	CO	N*	CH ₂	3	OEt
D15(c)	CH	N	CH	CH	-	CO	N*	CH ₂	3	OH
D16(a)	CH	CH	CH	N	CH	CH	CO	N*	3	OH
D16(b)	CH	CH	CH	N	CH ₂	CH ₂	CO	N*	3	OH
D17(a)	N	CH	CH	CH	CH	CH	CO	N*	3	OH
D17(b)	N	CH	CH	CH	CH ₂	CH ₂	CO	N*	3	OH
D20(d)	N	CH	CH	CH	O	CH ₂	CO	N*	2	OH
D22(b)	CH	CH	N	CH	O	CH ₂	CO	N*	2	OH

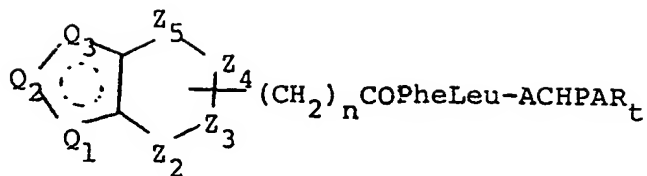
TABLE 3

Desc.No.	Q ₁	Q ₂	Q ₃	Z ₂	Z ₃	Z ₄	Z ₅	n	R _u
D18(a)	S	CH	CH	CO	N*	CH ₂	CH ₂	4	OEt
D18(b)	S	CH	CH	CO	N*	CH ₂	CH ₂	4	OH

TABLE 4

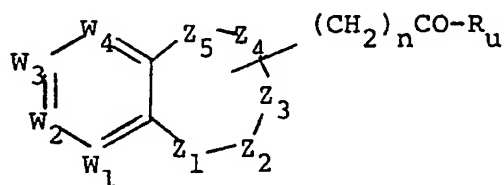
Ex. No.	W ₁	W ₂	W ₃	W ₄	Z ₂	Z ₃	Z ₄	Z ₅	n	R _t	salt/ solvate
E1	CH	CH	CH	N	-	O	CH ₂	CH*	1	NH ⁱ Bu	0.5H ₂ O
E2	CH	CH	CH	NO	-	O	CH ₂	CH*	1	NH ⁱ Bu	
E3	CH	CH	CH	N	O	CH ₂	CO	N*	2	API	0.5CHCl ₃
E4a	CH	CH	N	CH	S	CH ₂	CO	N*	2	API	2H ₂ O
E4b	CH	CH	N	CH	S	CH ₂	CO	N*	2	API	HCl
E5	CH	CH	CH	N	-	O	CH ₂	CH*	1	API	
E6	CH	CH	CH	N	-	NH	CO	N*	2	API	HCl
E7	N	CH	CH	CH	-	NH	CO	N*	2	API	HCl
E8	CH	CH	CH	NO	-	O	CH ₂	CH*	1	API	2HCl
E9	CH	CH	N	CH	SO ₂	CH ₂	CO	N*	2	API	HCl
E10	N	CH	CH	CH	S	CH ₂	CO	N*	2	API	HCl
E11	N	CH	CH	CH	SO ₂	CH ₂	CO	N*	2	API	HCl.1.5H ₂ O
E12	N	CH	CH	CH	-	CH ₂	N*	CO	3	API	HCl
E13	N	CH	CH	CH	-	CO	N*	CH ₂	3	API	HCl
E14	CH	N	CH	CH	-	CO	N*	CH ₂	3	API	2HCl
E15	CH	CH	CH	N	CH ₂	CH ₂	CO	N*	3	API	HCl
E16	N	CH	CH	CH	CH ₂	CH ₂	CO	N*	3	API	HCl
E19	N	CH	CH	CH	O	CH ₂	CO	N*	2	API	
E21	CH	CH	N	CH	O	CH ₂	CO	N*	2	API	

TABLE 5



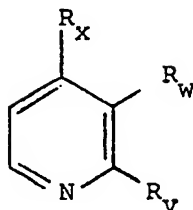
Ex.No.	Q ₁	Q ₂	Q ₃	Z ₂	Z ₃	Z ₄	Z ₅	n	R _t	Salt/ Solvate
E17a	S	CH	CH	CO	N*	CH ₂	CH ₂	4	API	H ₂ O
E17b	S	CH	CH	CO	N*	CH ₂	CH ₂	4	API	HCl.1.5H ₂ O
E21	S	CH	CH	CO	N*	CH ₂	CH ₂	3	API	

TABLE 6



Desc/ Ex.No.	W ₁	W ₂	W ₃	W ₄	Z ₁	Z ₂	Z ₃	Z ₄	Z ₅	n	R _u
D19(d)	N	CH	CH	CH	S	CH ₂	CH ₂	CO	N*	2	OMe
D19(e)	N	CH	CH	CH	S	CH ₂	CH ₂	CO	N*	2	OH
D21(c)	CH	CH	N	CH	S	CH ₂	CH ₂	CO	N*	2	OH
E18	N	CH	CH	CH	S	CH ₂	CH ₂	CO	N*	2	Phe-Leu-ACHPA-API
E20	CH	CH	N	CH	S	CH ₂	CH ₂	CO	N*	2	Phe-Leu-ACHPA-API

TABLE 7



Desc.No.	R _V	R _W	R _X
D7 (a)	NHCH ₂ CH ₂ CO ₂ Et	NO ₂	H
D7 (b)	NHCH ₂ CH ₂ CO ₂ Et	NH ₂	H
D8 (a)	NHCO ₂ Et	NHCO ₂ Et	H
D8 (b)	- NHCONH -		H
D11 (a)	- SCH ₂ CONH -		H
D18 (a)	SCH ₂ CH ₂ CO ₂ H	NO ₂	H
D19 (b)	SCH ₂ CH ₂ CO ₂ H	NH ₂	H
D19 (c)	- SCH ₂ CH ₂ CONH -		H
D20 (a)	OCH ₂ CO ₂ Et	NO ₂	H
D20 (b)	OCH ₂ CO ₂ Et	NH ₂	H
D20 (c)	- OCH ₂ CONH -		H
D21 (a)	H	NO ₂	SCH ₂ CH ₂ CO ₂ H
D21 (b)	H	- NHCOCH ₂ CH ₂ S -	
D22 (a)	H	- NHCOCH ₂ O -	

Description 1

2,3-Dihydrofuro[3,2-b]pyridine-3-acetic acid

10% Aqueous sodium hydroxide (0.7ml) was added dropwise to a stirred solution of ethyl 2,3-dihydrofuro[3,2-b]pyridine-3-acetate * (0.32g) in ethanol (5ml) at 0 °C, and the resultant mixture stirred at room temperature for 18h. Water (50ml) was added and the aqueous solution washed with chloroform. The aqueous layer was neutralised with 10% aqueous citric acid and evaporated in vacuo. The residue was triturated with chloroform and filtered and the filtrate was evaporated in vacuo to afford the title compound (0.17g) as a white solid.

NMR (δ) (DMSO_d₆): 2.1(1H,dd), 2.6(1H,dd), 3.65(1H,m), 4.25(1H,t), 4.8(1H,t), 7.05(2H,m), 7.95(1H,m)

* Tet Letts 1985, 26, 49, 6001-6004.

Description 2(a) Methyl 2,3-dihydro-3-oxo-4H-pyrido[3,2-b]-1,4-oxazine-4-propanoate

5

Methyl acrylate (0.18ml) was added dropwise to a stirred mixture of 2,3-dihydro-3-oxo-4H-pyrido[3,2-b]-1,4-oxazine (0.50g) and anhydrous potassium carbonate (0.41g) in dry dimethylformamide (DMF) (5ml). The reaction mixture was stirred at room temperature for 66h, poured into water and extracted with chloroform (3x100ml). Combined extracts were dried (Na_2SO_4) and the solvent removed in vacuo. The crude product
 10 was purified by chromatography on silica gel using ethyl acetate/petroleum ether (b.p. 60-80 °C) (0-30% ethyl acetate, gradient) to afford the title compound (0.48g) as a white solid.
 NMR (δ) (CDCl_3): 2.75(2H,t), 3.65(3H,s), 4.45(2H,t), 4.7(2H,s), 6.9(1H,dd), 7.2(1H,dd), 8.0(1H,dd).

15 (b) 2,3-Dihydro-3-oxo-4H-pyrido[3,2-b]-1,4-oxazine-4-propanoic acid

10% Aqueous sodium hydroxide (0.56ml) was added to a chilled, stirred solution of methyl 2,3-dihydro-3-oxo-4 H -pyrido[3,2-b]-1,4-oxazine-4-propanoate (0.47g) in methanol (5ml) and the resultant mixture stirred at room temperature for 21h. Solvents were evaporated in vacuo and the residue dissolved in water
 20 and adjusted to neutrality with 2M hydrochloric acid. The solution was freeze-dried and the residue triturated with methanol/dichloromethane (1:5 mixture). Filtration and evaporation of the filtrate afforded the title compound (0.40g) as a white solid.
 NMR (δ) (DMSO-d_6): 2.2(2H,t), 4.1(2H,t), 4.7(2H,s), 7.0(1H,m), 7.35(1H, dd), 8.0(1H,dd).

25

Description 3(a) BOC-ACHPA 3-(1-imidazolyl)propylamide

30

To a stirred ice cooled mixture of BOC-ACHPA-OH (1.2g) and 1-hydroxybenzotriazole (HOBT) (0.51g) in dry DMF (10 ml) was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide.HCl (DEC) (0.73g). After 10 min N-(3-aminopropyl)imidazole (0.68ml) was added and stirring continued for 16h. Water was added and the solution extracted with ethyl acetate (3x200ml). The combined extracts were washed with water, saturated
 35 sodium hydrogen carbonate and brine, dried (Na_2SO_4) and the solvent evaporated in vacuo. The residue was purified by chromatography on silica gel using methanol/chloroform (0-3% methanol, gradient) to afford the title compound (1.45g).
 NMR (δ) (DMSO-d_6): 0.7-1.9(24H,m), 2.15(2H,m), 3.0(2H,m), 3.8(1H,m), 3.95(2H,t), 4.7(1H,m), 6.25(1H,d), 7.1-(1H,s), 7.6(1H,s) 7.8(1H,t).

40

(b) CBZ-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide

Chilled trifluoroacetic acid (TFA) (20ml) was added to a stirred solution of BOC-ACHPA 3-(1-imidazolyl)-propylamide (1.4g) in dry dichloromethane (20ml) and the reaction mixture stirred at room temperature for
 45 1h. Evaporation in vacuo afforded a light yellow foam (1-8g). This material was dissolved in dry DMF (10 ml) and triethylamine (1.37ml) and added to a stirred mixture of CBZ-Phe-Leu-OH (2.02g), HOBT.(0.66g) and DEC (0.94g) in DMF (20ml) at 0 °C. After 16h water was added and the mixture extracted with ethyl acetate (3x250ml). Combined extracts were washed with water, saturated sodium hydrogen carbonate and brine,
 50 dried (Na_2SO_4) and evaporated in vacuo. The crude product was purified by chromatography on silica gel using methanol/chloroform (0-5% methanol, gradient) to afford the title compound (1.95g).
 NMR (δ) (CDCl_3) 0.7-2.5(26H,m), 2.8-3.4(6H,m), 3.9-4.5(6H,m), 5.0(2H,m), 6.6(1H,m), 6.9-7.4(13H,m).
 M.S. (m/z) (FAB) (M + 1) = 717 (consistent with m.w. = 716)

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(c) Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide

CBZ-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide (1.8g) in ethanol (150ml) was hydrogenated at room

temperature and pressure in the presence of 10% palladium on carbon for 20h. Filtration through kieselguhr and evaporation of the filtrate afforded the crude product which was purified by chromatography on silica gel using methanol/chloroform (0-12% methanol, gradient) to afford the title compound (0.31g).

NMR (δ) (CD_3OD): 0.8-1.7(22H,m), 2.0(2H,m), 2.25(2H,m), 2.8(1H,dd) 3.0-3.2(3H,m), 3.65(1H,m), 3.9-4.2-
 5 (4H,m), 4.3(1H,m), 6.9(1H,s), 7.15(1H,s), 7.25(5H,m), 7.7(1H,s).

Description 4

10

(a) BOC-Phe-Leu-ACHPA isobutylamide

This material was formed from BOC-Phe-Leu-OH (1.48g) and ACHPA isobutylamide * (1.06g), following the procedure of Description 3(a). The crude product was chromatographed on silica gel using
 15 methanol/chloroform (0-2% methanol, gradient). This gave the title compound (1.91g).

NMR (δ) (CDCl_3): 0.7-2.3 (31 H,m), 2.8-3.2(4H,m), 3.8-4.3(4H,m), 6.4-7.4(8H,m).

(b) Phe-Leu-ACHPA isobutylamide.TFA

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BOC-Phe-Leu-ACHPA isobutylamide (2.32g) was dissolved with stirring in TFA (10ml) at 0° C. After 1h, solvent was removed in vacuo, and the residue was triturated with ether to give the title compound.

NMR (δ) (CDCl_3): 0.6-2.5 (31H,m), 2.8-3.3(4H,m), 3.7-4.4(4H,m), 7.0-8.0(8H,m).

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Description 5

(a) CBZ-Phe-Leu-ACHPA isobutylamide

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This material was formed from CBZ-Phe-Leu-OH (4.13g), following the procedure of Description 4(a). The crude product was purified by chromatography on silica gel using methanol/chloroform (0-3% methanol, gradient). This gave the title compound (6.40g) as a light brown solid.

NMR (δ) (CDCl_3): 0.75-1.85 (29H,m), 2.3(2H,m), 2.9-3.15 (4H,m), 4.0(2H,m), 4.3-4.5(2H,m), 5.05(2H,s), 5.5-
 35 (1H,d), 6.5-6.85(3H,m), 7.1-7.4(11H,m).

(b) Phe-Leu-ACHPA isobutylamide.CH₃CO₂H

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This material was formed from CBZ-Phe-Leu-ACHPA isobutylamide (6.4g) following the procedure of Description 3(c), but using ethanol/acetic acid as solvent. This gave the title compound (7.4g), still containing residual acetic acid.

NMR (δ) (CDCl_3): 0.8-1.85(29H,m), 2.05(m), 2.35(2H,m), 2.9-3.2(4H,m), 3.95(2H,m), 4.15(1H,m), 4.3(1H,m),
 45 7.25(5H,m), 8.0(b).

(c) Phe-Leu-ACHPA isobutylamide

Phe-Leu-ACHPA isobutylamide.CH₃CO₂H (7.4g) was dissolved in ethyl acetate and extracted into
 50 aqueous citric acid. The extract was basified with sodium carbonate, and the product was isolated by extraction into chloroform, drying (Na_2SO_4) and evaporation in vacuo. This gave the title compound (3.2g).

NMR (δ) (CDCl_3): 0.7-1.0 (13H,m), 1.0-1.9(16H,m), 2.3(2H,ABX), 2.75(1H,dd), 3.05(2H,m), 3.2(1H,dd), 3.65-
 (1H,dd), 3.95(2H,m), 4.35(1H,m), 6.5(1H,d), 6.6(1H,t), 7.15-7.35(6H,m), 7.7(1H,d).

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Description 6

* J. Med. Chem. 1985, 28, 1779-1790.

a) Methyl 2,3-dihydro-3-oxo-4H-pyrido[4,3-b]-1,4-thiazine-4-propanoate

This material was prepared from 2,3-dihydro-3-oxo-4 H -pyrido[4,3-b]-1,4-thiazine * (0.18g), following the procedure of Description 2(a). Purification by chromatography on silica gel using methanol/chloroform (0-4% methanol, gradient) afforded the title compound (0.18g) as a colourless waxy solid.

NMR (δ) (CDCl_3): 2.75 (2H,t), 3.45 (2H,s), 3.7 (3H,s), 4.35 (2H,t), 7.3 (1H,dd), 8.2 (1H,d), 8.45 (1H,s).

b) 2,3-Dihydro-3-oxo-4H-pyrido[4,3-b]-1,4-thiazine-4-propanoic acid

This material was prepared from methyl 2,3-dihydro-3-oxo-4 H -pyrido[4,3-b]-1,4-thiazine-4-propanoate (0.17g), following the procedure of Description 2(b). This gave the title compound (0.15g) as a white solid.

NMR (δ) (DMSO-d_6): 2.15 (2H,m), 3.55 (2H,s), 4.1 (2H,m), 7.4 (1H,d), 8.1 (1H,d), 8.5 (1H,s).

Description 7a) Ethyl 3-((3-nitro-2-pyridyl)amino)propanoate

2-Chloro-3-nitropyridine (2.0g), ethyl 3-amino-propanoate.HCl (1.94g) and anhydrous potassium carbonate (3.48g) were stirred in DMF (20 ml) for 16h. The mixture was evaporated in vacuo, partitioned between chloroform and water, and separated. The aqueous portion was extracted with chloroform, and the combined organics were washed with water and brine, dried (Na_2SO_4) and evaporated in vacuo. Chromatography on silica gel using methanol/chloroform (0-2% methanol, gradient) gave the title compound (2.5g).

NMR (δ) (CDCl_3): 1.3 (3H,t), 2.7 (2H,t), 3.95 (2H,q), 4.2 (2H,q), 6.7 (1H,dd), 8.4-9.7 (2H,m).

b) Ethyl 3-((3-amino-2-pyridyl)amino)propanoate

This material was formed from ethyl 3-((3-nitro-2-pyridyl)amino)propanoate (2.90g), following the procedure of Description 3(c). This gave the title compound (2.43g) as a red oil.

NMR (δ) (CDCl_3): 1.3 (3H,t), 2.7 (2H,t), 3.2 (2H,b), 3.75 (2H,m), 4.15 (2H,q), 4.8 (1H,b), 6.5 (1H,dd), 6.85 (1H,d), 7.7 (1H,d).

c) Ethyl 2,3-dihydro-2-oxo-1H-imidazo[4,5-b]-e-3-propanoate

Ethyl 3-((3-amino-2-pyridyl)amino)propanoate (2.4g) was stirred in dichloromethane (50 ml) as 1,1'-carbonyldiimidazole (2.27g) was added. After a vigorous reaction, the mixture was stirred for 16h, evaporated in vacuo, dissolved in ethyl acetate, washed with water, dried (Na_2SO_4) and evaporated in vacuo. Chromatography on silica gel using methanol/chloroform (0-5% methanol, gradient) gave the title compound (2.22g) as a purple solid.

NMR (δ) (CDCl_3): 1.2 (3H,t), 2.9 (2H,t), 4.15 (2H,q), 4.35 (2H,t), 5.7 (b), 7.0 (1H,dd), 7.3 (1H,dd), 8.05 (1H,dd), 10.4 (b).

d) 2,3-Dihydro-2-oxo-1H-imidazo[4,5-b]pyridine-3-propanoic acid

This material was formed from ethyl 2,3-dihydro-2-oxo-1 H -imidazo[4,5-b]pyridine-3-propanoate (0.82g), following the procedure of Description 2(b), but using ethanol as solvent. This gave the title compound (0.8g) as a pink solid.

NMR (δ) (DMSO-d_6): 2.3 (2H,t), 3.95 (2H,t), 6.9 (1H,dd), 7.25 (1H,d), 7.85 (1H,d).

* Synth. Comm. 331-3 (1985).

Description 8a) 2,3-Di(ethoxycarbonylamino)pyridine

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2,3-Diaminopyridine (2.77g) and dry pyridine (8.1 ml) were stirred at 0 °C in dry tetrahydrofuran (THF) (50 ml) as ethyl chloroformate (4.9 ml) was added dropwise. The mixture was stirred at 0 °C for 15 min, and then allowed to rise to ambient temperature over 3h. It was diluted with ethyl acetate, washed with water, dried (Na₂SO₄) and evaporated in vacuo. Chromatography on alumina using 10% ethyl acetate in petroleum ether (b.p. 60-80 °C) gave the title compound (3.16g) as an orange solid.

NMR (δ) (CDCl₃): 1.35 (6H, 2xt), 4.25 (4H, 2xg), 7.2 (1H,dd), 8.1 (1H,d), 8.2 (1H,d), 8.45 (1H,b), 8.8 (b).

b) 2,3-Dihydro-2-oxo-1H-imidazo[4,5-b]pyridine

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2,3-Di(ethoxycarbonylamino)pyridine (3.16g) was dissolved in 2M aqueous sodium hydroxide (100 ml) at 50 °C. The solution was then acidified (5M hydrochloric acid) and neutralised with sodium hydrogen carbonate, and extracted with ethyl acetate. The extract was dried (Na₂SO₄) and evaporated in vacuo to give the title compound (0.20g), as a brown solid. On standing, the aqueous portion yielded a brown precipitate, which was filtered off and dried in vacuo, affording a further crop of the title compound (1.14g).

NMR (δ) (DMSO-d₆): 6.9 (1H,dd), 7.2 (1H,d), 7.8 (1H,d), 11.1 (1H,vb).

c) Methyl 2,3-dihydro-2-oxo-1H-imidazo[4,5-b]-pyridine-1-propanoate

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This material was formed from 2,3-dihydro-2-oxo-1H-imidazo[4,5-b]pyridine (1.3g), following the procedure of Description 2(a). The crude product was purified by chromatography using ethyl acetate/petroleum ether (b.p. 60-80 °C) (0-100% ethyl acetate, gradient). This gave the title compound (1.2g).

NMR (δ) (CDCl₃): 2.85 (2H,t), 3.65 (3H,s), 4.2 (2H,t), 7.05 (1H,dd), 7.4 (1H,d), 8.1 (1H,d).

d) 2,3-Dihydro-2-oxo-1H-imidazo[4,5-b]pyridine-1-propanoic acid

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This material was formed from methyl 2,3-dihydro-2-oxo-1H-imidazo[4,5-b]pyridine-1-propanoate (0.78g), following the procedure of Description 2(b). This gave the title compound (0.42g) as a white solid.

NMR (δ) (DMSO-d₆): 2.25 (2H,t), 3.95 (2H,t), 6.9 (1H,dd), 7.4 (1H,d), 7.8 (1H,d).

40 Description 9a) BOC-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide

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This material was formed from BOC-ACHPA 3-(1-imidazolyl)propylamide (1.53g) (Description 3(a)) and BOC-Phe-Leu-OH (1.60g), following the procedure of Description 3(b). The crude product was chromatographed on silica gel using methanol/chloroform (0-10% methanol, gradient). This gave the title compound (1.95g).

NMR (δ) (CDCl₃): 0.7-1.05 (8H,m), 1.05-1.9 (2H,m), 2.0 (2H,m), 2.2-2.5 (2H,m), 2.9-3.6 (5H,m), 3.9-4.5 (6H,m), 5.1 (b), 6.4 (b), 6.6 (bd), 7.0 (1H,s), 7.05 (1H,s), 7.15-7.4 (m), 7.65 (1H,s).

b) Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide .2(CF₃CO₂H)

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To a solution of BOC-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide (0.60g) in dichloromethane (6 ml) was added trifluoroacetic acid (6 ml). The mixture was stirred for 1h, and evaporated. Trituration with ether/pentane and filtration then gave the title compound (0.62g) as a white solid.

Description 10

a) t-Butyl 2,3-dihydro-3-oxo-4H-pyrido[4,3-b]-1,4-thiazine-4-propanoate

This material was formed from 2,3-dihydro-3-oxo-4 H -pyrido[4,3-b]-1,4-thiazine (0.36g) and t-butyl acrylate (0.63 ml), following the procedure of Description 2(a). This gave the title compound (0.62g) as a pale orange oil.

NMR (δ) (CDCl_3): 1.4 (9H,s), 2.65 (2H,t), 3.45 (2H,s), 4.3 (2H,t), 7.3 (1H,d), 8.2 (1H,d), 8.45 (1H,s).

b) t-Butyl 2,3-dihydro-1,1,3-trioxo-4H-pyrido[4,3-b]-1,4-thiazine-4-propanoate

t-Butyl 2,3-dihydro-3-oxo-4H-pyrido[4,3-b]-1,4-thiazine-4-propanoate (0.18g) and monoperoxyphthalic acid, magnesium salt .6H₂O (0.60g) were stirred in methanol (5 ml) for 16h. The mixture was evaporated in vacuo, taken up in ether, washed with 5% sodium hydrogen carbonate solution and brine, dried (Na_2SO_4) and evaporated in vacuo. The residue was taken up in chloroform, filtered and evaporated in vacuo, giving the title compound (0.092g) as a colourless oil.

NMR (δ) (CDCl_3): 1.4 (9H,s), 2.7 (2H,t), 4.3 (2H,s), 4.4 (2H,t), 7.8 (1H,d), 8.7 (1H,d), 8.9 (1H,s).

c) 2,3-Dihydro-1,1,3-trioxo-4H-pyrido[4,3-b]-1,4-thiazine-4-propanoic acid

t-Butyl 2,3-dihydro-1,1,3-trioxo-4 H -pyrido[4,3-b]-1,4-thiazine-4-propanoate (0.15g) was dissolved in dichloromethane (2 ml) and TFA (2 ml), and left to stand for 3h. Solvent was evaporated in vacuo, and the residue was triturated with Et₂O. Filtration then gave the title compound (0.063g) as a white powder.

NMR (δ) (DMSO-d_6): 2.6 (2H,t), 4.3 (2H,t), 4.95 (2H,s), 7.85 (1H,d), 8.65 (1H,d), 9.05 (1H,s).

Description 11

a) 2,3-Dihydro-3-oxo-4H-pyrido[2,3-b]-1,4-thiazine

2-Chloro-3-nitropyridine (2.31g), anhydrous potassium carbonate (4.02g) and ethyl mercaptoacetate (2.4 ml) was stirred in DMF (30 ml) for 1.5h. The mixture was diluted with ethyl acetate (300 ml), washed with water and brine, dried (Na_2SO_4) and evaporated in vacuo to give a dark green mobile oil (4.53g). This was dissolved in glacial acetic acid (100 ml), and stirred as iron powder (20g) was added. The mixture was stirred at 90 °C for 2h, cooled, and evaporated in vacuo. The resultant brown solid was extracted twice with xylene, and the extracts were evaporated to give the title compound (0.85g), impure, as a brown solid. The residue remaining after xylene extraction was dissolved in 10% aq. acetic acid, filtered, and extracted with 10% methanol/chloroform. This extract was dried (Na_2SO_4) and evaporated in vacuo, giving a second crop of material (0.88g), pure, as a buff solid. The initial xylene extract was purified by trituration with ether, giving a total yield of 1.38g.

NMR (δ) (DMSO-d_6): 3.65 (2H,s), 7.2 (2H,m), 8.1 (1H,dd), 10.65 (1H,b).

b) t-Butyl 2,3-dihydro-3-oxo-4H-pyrido[2,3-b]-1,4-thiazine-4-propanoate

This material was formed from 2,3-dihydro-3-oxo-4 H -pyrido[2,3-b]-1,4-thiazine (0.84g), following the procedure of Description 10(a). This gave the title compound (1.57g) as a brown oil.

NMR (δ) (CDCl_3): 1.4 (9H,s), 2.6 (2H,t), 3.5 (2H,s), 4.2 (2H,t), 7.2 (1H,dd), 7.5 (1H,dd), 8.2 (1H,dd).

c) 2,3-Dihydro-3-oxo-4H-pyrido[2,3-b]-1,4-thiazine-4-propanoic acid

This material was formed from t-butyl 2,3-dihydro-3-oxo-4 H -pyrido[2,3-b]-1,4-thiazine-4-propanoate (0.33g), following the procedure of Description 10(c). This gave the title compound (0.24g) as a white

powder.

NMR (δ) ($\text{CDCl}_3/\text{DMSO-d}_6$): 2.55 (m), 3.5 (s), 4.15 (2H,t), 7.25 (1H,dd), 7.65 (1H,d), 8.1 (1H,dd).

5 Description 12

a) t-Butyl 2,3-dihydro-1,1,3-trioxo-4H-pyrido[2,3-b]-1,4-thiazine-4-propanoate

10 This material was formed from t-butyl 2,3-dihydro-3-oxo-4 H -pyrido[2,3-b]-1,4-thiazine-4-propanoate (Description 11(b)) (0.40g), following the procedure of Description 10(b). This gave the title compound (0.29g) as a clear oil, but contaminated with ca.20% of an unidentified impurity (NMR).

NMR (δ) (CDCl_3): 1.4 (9H,s), 2.65 (2H,t), 4.3 (2H,t), 4.35 (2H,s), 7.65 (1H,dd), 7.9 (1H,d), 8.55 (1H,dd).

15

b) 2,3-Dihydro-1,1,3-trioxo-4H-pyrido[2,3-b]-1,4-thiazine-4-propanoic acid

This material was formed from t-butyl 2,3-dihydro-1,1,3-trioxo-4 H -pyrido[2,3-b]-1,4-thiazine-4-propanoate (0.26g), following the procedure of Description 10(c). This gave the pure title compound (0.16g) as a grey solid.

20

NMR (δ) (DMSO-d_6): 2.55 (2H,t), 4.2 (2H,t), 4.95 (2H,s), 7.8 (1H,dd), 8.2 (1H,d), 8.5 (1H,d).

Description 13

25

a) Ethyl 5,7(6H)-dioxo-5H-pyrrolo[3,4-b]pyridine-6-butanoate

A mixture of pyridine-2,3-dicarboxylic acid (15g) and urea (6.65g) was heated at 180 °C for 30 min. After cooling to room temperature, the residue was triturated with ethyl acetate and water to give the crude dicarboximide intermediate.

30

This crude intermediate was alkylated with ethyl 4-bromobutyrate following the procedure of Description 2(a) to give the title compound (3.75g).

NMR (δ) (CDCl_3): 1.3 (3H,t), 2.0 (2H,quin), 2.4 (2H,t), 3.8 (2H,t), 4.1 (2H,q), 7.6 (1H,dd), 8.1 (1H,d), 9.0 (1H,d).

35

b) Ethyl 6,7-dihydro-7-hydroxy-5-oxo-5H-pyrrolo-[3,4-b]pyridine-6-butanoate

To a stirred solution of ethyl 5,7(6H)-dioxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoate (0.95g) in methanol (55ml) at 0 °C was added sodium borohydride (0.28g). After 20 min, the reaction was quenched by the addition of 5M hydrochloric acid, and the reaction mixture evaporated. The residue was partitioned between chloroform (50 ml) and aqueous sodium hydrogen carbonate (20 ml), then the organic phase was dried and evaporated. Chromatography on silica using ethyl acetate gave the title compound (0.68g).

40

NMR (δ) (CDCl_3): 1.2 (3H,t), 2.0-2.2 (2H,m), 2.4-2.5 (2H,m), 3.6-3.7 (2H,m), 4.1 (2H,q), 7.5 (1H,dd), 8.1 (1H,d), 8.8 (1H,d).

45

c) Ethyl 6,7-dihydro-5-oxo-5H-pyrrolo[3,4-b]-pyridine-6-butanoate

A solution of ethyl 6,7-dihydro-7-hydroxy-5-oxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoate (0.33g) in acetic acid (8 ml) was maintained under reflux for 13h in the presence of zinc dust (0.42g). The reaction mixture was filtered and evaporated, then the title compound (0.24g) was obtained by chromatography on silica using ethyl acetate.

50

NMR (δ) (CDCl_3): 1.2 (3H,t), 2.0 (2H, quin), 2.4 (2H,t), 3.7 (2H,t), 4.0 (2H,q), 4.8 (2H,s), 7.7 (1H,t), 8.4 (1H,d), 8.9 (1H,bs).

55

d) 6,7-Dihydro-5-oxo-5H-pyrrolo[3,4-]pyridine-6-butanoic acid

To a solution of ethyl 6,7-dihydro-5-oxo-5H-pyrrolo[3,4-b]pyridine-6-butanoate (0.36g) in ethanol (7 ml) was added water (7 ml) and potassium carbonate (0.20g). This mixture was stirred at room temperature for 3 days, then evaporated. Chromatography on silica using 50% methanol/chloroform gave the title compound (0.18g).

NMR (δ) (DMSO $_d$): 1.8-1.9 (2H,quin), 2.25 (2H,t), 3.5 (2H,t), 4.5 (2H,s), 7.5 (1H,m), 8.1 (d), 8.7 (1H,d).

10 Description 14a) Ethyl 6,7-dihydro-5-hydroxy-7-oxo-5H-pyrrolo-[3,4-b]pyridine-6-butanoate

15 The title compound (0.23g) was obtained as a more slowly eluting by-product of the synthesis of ethyl 6,7-dihydro-7-hydroxy-5-oxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoate (Description 13(b)).

NMR (δ) (CDCl $_3$): 1.3 (3H,t), 1.9-2.1 (2H,m), 2.4-2.5 (2H,m), 3.5-3.8(2H,m), 4.0 (2H,q), 7.4-7.5 (1H,m), 8.0 (1H,d), 8.7 (1H,d).

20

b) Ethyl 6,7-dihydro-7-oxo-5H-pyrrolo[3,4-b]pyridine-6-butanoate

A solution of ethyl 6,7-dihydro-5-hydroxy-7-oxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoate (0.23g) in ethanol (25 ml) was acidified with 5M hydrochloric acid, then hydrogenated in the presence of 10% Pd/C for 4 days. The mixture was filtered and the filtrate evaporated to give the title compound (0.17g).

NMR (δ) (DMSO $_d$): 1.1-1.2 (3H,m), 1.8-1.9 (2H,m), 2.3 (2H,t), 3.6 (2H,t), 4.0 (2H,q), 4.5 (2H,s), 7.6 (1H,dd), 8.1 (1H,d), 8.7 (1H,d).

30 c) 6,7-Dihydro-7-oxo-5H-pyrrolo[3,4-b]pyridine-6-butanoic acid

This material was formed from ethyl 6,7-dihydro-7-oxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoate (0.17g), following the procedure of Description 13(d). This gave the title compound (0.12g).

NMR (δ) (DMSO $_d$): 1.8-1.9 (2H,m), 2.2-2.3 (2H,m), 3.6 (2H,t), 4.5 (2H,s), 7.6 (1H,dd), 8.1 (1H,d), 8.7(1H,d).

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Description 1540 a) Ethyl 5,7(6H)-dioxo-5H-pyrrolo[3,4-c]pyridine-6-butanoate

This material was formed from pyridine-3,4-dicarboximide (2.5g) and ethyl 4-bromobutyrate (2.0ml), following the procedure of Description 2(a). This gave the title compound (3.13g).

NMR (δ) (CDCl $_3$): 1.25 (3H,t), 2.0 (2H, quin), 2.4 (2H,t), 3.8 (2H,t), 4.1 (2H,q), 7.8 (1H,d), 9.05 (1H,d), 9.15 (1H,s).

45

b) Ethyl 6,7-dihydro-5H-7-oxopyrrolo[3,4-c]pyridine-6-butanoate

50 This material was formed from ethyl 5,7(6H)-dioxo-5 H -pyrrolo[3,4-c]pyridine-6-butanoate (0.90g), following the procedures of Descriptions 13(b) and 13(c), but omitting chromatography of the 5-hydroxy intermediate. Chromatography on silica using ethyl acetate gave the title compound (0.41g).

NMR (δ) (CDCl $_3$): 1.2 (3H,t), 2.0 (2H, quin), 2.4 (2H,t), 3.7 (2H,t), 4.05 (2H,q), 4.45 (2H,s), 7.45 (1H,d), 8.75 (1H,d), 9.1 (1H,s).

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c) 6,7-Dihydro-5H-7-oxopyrrolo[3,4-c]pyridine-6-butanoic acid.HCl

This material was formed from ethyl 6,7-dihydro-5 H -7- oxopyrrolo[3,4-c]pyridine-6-butanoate (0.30g), following the procedure of Description 13(d), followed by acidification/freeze-drying by the procedure of Example 4(b). This gave the title compound (0.20g).

NMR (δ) (DMSO $_d$): 1.8 (2H, quin), 2.25 (2H,t), 3.55 (2H,t), 4.7(2H,s), 5.0-6.5 (3H,b), 8.1 (1H,d), 8.9 (1H,d), 9.15 (1H,s).

Description 16

10

a) 7,8-Dihydro-7-oxo-8H-1,8-naphthyridine-8-butanoic acid

A mixture of 1,8-naphthyridine * (2.5g) and ethyl 4-bromobutyrate (13.7 ml) in ethanol (100 ml) was stirred at reflux for 6 days, then evaporated in vacuo. Water (50 ml) was added and the mixture stirred in an ice bath as a solution of potassium ferricyanide (38g) in water (125 ml) was added dropwise over 0.5h, followed by the dropwise addition of a solution of potassium hydroxide (17.5g) in water (25 ml). Stirring was continued for 24h, during which time the reaction mixture was allowed to warm to room temperature. Water was added to dissolve the remaining solid and the solution washed with chloroform. The aqueous layer was acidified with 5M hydrochloric acid and extracted with chloroform. Pooled chloroform extracts were dried (Na $_2$ SO $_4$) and evaporated in vacuo. The residue was recrystallised from methanol to afford the title compound (2.0g).

NMR (δ) (DMSO $_d$): 1.9 (2H,m), 2.2 (2H,m), 4.4 (2H,m), 6.7 (1H,d), 7.3 (1H,dd), 7.9(1H,d), 8.2 (1H,dd), 8.6 (1H,m), 12.05 (1H,b).

25

b) 7-Oxo-5,6,7,8-tetrahydro-8H-1,8-naphthyridine-8-butanoic acid

To stirred palladium black (0.2g) under nitrogen was added 5% formic acid/methanol (8 ml), followed by a solution of 7,8-dihydro-7-oxo-8 H -1,8-naphthyridine-8-butanoic acid (0.40g) in 5% formic acid/methanol (24 ml) in two portions, the latter after 1h reaction. The mixture was stirred at room temperature overnight, filtered through celite and the filtrate evaporated in vacuo. The residue was purified by chromatography on silica gel using methanol/chloroform (0-20% methanol, gradient) to afford the title compound (0.06g).

NMR (δ) (CDCl $_3$): 2.0 (2H,m), 2.4 (2H,t), 2.7 (2H,t), 2.9 (2H,t), 4.2 (2H,t), 6.95 (1H,m), 7.45 (1H,d), 8.2 (1H,d), 9.8 (1H,b).

35

Description 17

40 a) 5,6-Dihydro-6-oxo-5H-1,5-naphthyridine-5-butanoic acid

This material was formed from 1,5-naphthyridine * (1.00g) following the procedure of Description 16(a). On evaporation of the chloroform extracts, the solid produced was filtered from the-residual liquid to afford the title compound (0.12g).

45 NMR (δ) (DMSO $_d$): 1.8 (2H,m), 2.35 (2H,t), 4.2 (2H,t), 6.9 (1H,dd), 7.6 (1H,dd), 7.9 (1H,dd), 8.1 (1H,d), 9.45 (1H,d), 12.1 (1H,b).

b) 6-Oxo-5,6,7,8-tetrahydro-5H-1,5-naphthyridine-5-butanoic acid

50

This material was formed from 5,6-dihydro-6-oxo-5 H -1,5-naphthyridine-5-butanoic acid (0.12g), following the procedure of Description 16(b). Isolation by filtration through celite and evaporation of the filtrate in vacuo afforded the title compound (0.12g) which was used without further purification.

55 NMR (δ) (CD $_3$ OD): 1.9 (2H,m), 2.4 (2H,t), 2.75 (2H,t), 3.1 (2H,t), 4.0 (2H,t), 7.35 (1H,dd), 7.7 (1H,d), 8.1 (1H,d).

* Chem. Pharm. Bull. 19 (9) 1857-62 (1971).

Description 18a) Ethyl 7-oxo-4,5,6,7-tetrahydro-6H-thieno[2,3-c]-pyridine-6-pentanoate

5

This material was formed from 4,5-dihydrothieno[2,3-c]-pyrid-7(6H)-one * (0.33g) and ethyl 5-bromovalerate (0.34 ml) following the procedure of Description 2(a). The crude product was purified by chromatography on silica gel using ethyl acetate/petroleum ether (b.p. 60-80 °C) (0-30% ethyl acetate, gradient) to give the title compound (0.35g).

10 NMR (δ) (CDCl_3): 1.2 (3H,t), 1.7 (3H,m), 2.3 (2H,m), 2.9 (2H,t), 3.4-3.7 (4H,m), 4.1 (2H,q), 6.9 (1H,d), 7.4 (1H,d).

b) 7-Oxo-4,5,6,7-tetrahydro-6H-thieno[2,3-c]-pyridine-6-pentanoic acid

15

This material was formed from ethyl 7-oxo-4,5,6,7-tetrahydro-6H-thieno[2,3-c]pyridine-6-pentanoate (0.32g) following the procedure of Description 7(d). The title compound (0.27g) was isolated as a slightly coloured oil.

NMR (δ) (DMSO-d_6): 1.5 (3H,m), 2.2 (2H,m), 2.9 (2H,t), 3.4-3.5 (3H,m), 3.6 (3H,t), 7.0 (1H,d), 7.7 (1H,d).

20

Description 1925 a) 3-((3-Nitro-2-pyridyl)thio)propanoic acid

3-Mercaptopropionic acid (5.0 ml), 2-chloro-3-nitropyridine (9.11g) and anhyd. potassium carbonate (19.8g) were stirred in DMF (100 ml) for 5.5h. Solids were then dissolved by addition of water (200 ml), and the mixture was stirred for 16h, diluted with water (500 ml), washed with ethyl acetate, and acidified with

30

conc. hydrochloric acid. The mixture was then extracted into ethyl acetate, and the extract was dried

(Na_2SO_4) and evaporated in vacuo, giving the title compound (13.10g) as a yellow solid.

NMR (δ) ($\text{CDCl}_3/\text{DMSO-d}_6$): 2.7 (2H,t), 3.4 (2H,t), 7.3 (1H,dd), 8.55 (1H,dd), 8.8 (1H,m).

35 b) 3-((3-Amino-2-pyridyl)thio)propanoic acid

3-((3-Nitro-2-pyridyl)thio)propanoic acid (9.28g) was hydrogenated in ethanol (200 ml) over 10% palladium on charcoal (60% aq.paste, 4.0g) for 24h. A further 4.1g catalyst was then added, and hydrogenation continued for a further 64h. Filtration through kieselguhr, and evaporation in vacuo then gave

40

the title compound (7.54g) as an orange-yellow solid.

NMR (δ) (DMSO-d_6): 2.6 (2H,t), 3.3 (2H,t), 5.0 (2H,b), 6.9 (2H,m), 7.75 (1H,m).

c) 3,4-Dihydro-4-oxo-2H-pyrido[2,3-b]-1,4-thiazepine

45

3-((3-Amino-2-pyridyl)thio)propanoic acid (7.54g) and 1,3-dicyclohexylcarbodiimide (7.84g) were stirred together in DMF (75 ml) for 4.5h, evaporated in vacuo, taken up in ethyl acetate, filtered, and evaporated to a yellow/brown solid. This was purified by chromatography on silica gel using ethyl acetate/petroleum ether (b.p. 60-80 °C) (50-100% ethyl acetate, gradient), giving the title compound (1.00g).

50

NMR (δ) (CDCl_3): 2.7 (2H,t), 3.6 (2H,t), 7.3 (1H,dd), 7.4 (1H,dd), 8.05 (1H,b), 8.4 (1H,dd).

d) Methyl 3,4-dihydro-4-oxo-2H-pyrido[2,3-b]-1,4-thiazepine-5-propanoate

55

This material was formed from 3,4-dihydro-4-oxo-2H-pyrido[2,3-b]-1,4-thiazepine (0.55g) following the procedure of Description 2(a). This gave the title compound.

* J. Het. Chem. 3, 466 (1966).

NMR (δ) (CDCl_3): 2.6 (2H,t), 2.75 (2H,t), 3.4-3.6 (5H,m), 7.4 (1H,dd), 7.65 (1H,dd), 8.5 (1H,dd).

e) 3,4-Dihydro-4-oxo-2H-pyrido[2,3-b]-1,4-thiazepine-5-propanoic acid

5

This material is formed from methyl 3,4-dihydro-4-oxo-2H-pyrido[2,3-b]-1,4-thiazepine-5-propanoate, following the procedure of Description 7(d).

10 Description 20

a) Ethyl (3-nitro-2-pyridyl)oxyacetate

15

Sodium hydride (80% dispersion in mineral oil, 0.31g) was stirred under nitrogen in DMF (10 ml) as ethyl glycolate (1.0 ml) was added dropwise. After 10 min, 2-chloro-3-nitropyridine (1.65g) was added. After a further 15 min, the mixture was partitioned between ethyl acetate and water, and separated. The organic portion was washed with water and brine, dried (Na_2SO_4) and evaporated in vacuo, giving the title compound (2.1g) as a brown oil. ^1H NMR revealed ca. 20% contamination by the pyridine starting material (peaks not quoted below).

20

NMR (δ) (CDCl_3): 1.3 (3H,t), 4.25 (2H,q), 5.05 (2H,s), 7.1 (1H,m), 8.35 (2H,m).

b) Ethyl (3-amino-2-pyridyl)oxyacetate

25

This material was formed from ethyl (3-nitro-2-pyridyl) oxyacetate (2.09g), following the procedure of Description 3(c). This gave the title compound (1.76g) as a yellow oil.

NMR (δ) (CDCl_3): 1.3 (3H,t), 4.2 (2H,q), 4.9 (2H,s), 6.75 (1H,dd), 6.95 (1H,dd), 7.5 (1H,dd).

30

c) 2,3-Dihydro-3-oxo-4H-pyrido[2,3-b]-1,4-oxazine

Ethyl (3-amino-2-pyridyl)oxyacetate (1.75g) was stirred at reflux in dry xylene (50 ml) for 16h, evaporated to dryness, taken up in 10% methanol/chloroform and filtered. The filtrate was concentrated, and purified by chromatography on silica gel using methanol/chloroform (0-10% methanol, gradient). This gave the title compound (0.78g) as a brown solid.

35

NMR (δ) ($\text{DMSO}-d_6$): 4.75 (2H,s), 7.0 (1H,ddd), 7.2 (1H,m), 7.8 (1H,m), 10.85 (1H,b).

40 d) 2,3-Dihydro-3-oxo-4H-pyrido[2,3-b]-1,4-oxazine-4-propanoic acid

This material is formed from 2,3-dihydro-3-oxo-4H-pyrido[2,3-b]-1,4-oxazine, following the procedures of Descriptions 2(a) and 7(d).

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Description 21

a) 3-((3-Nitro-4-pyridyl)thio)propanoic acid

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This material was formed from 4-chloro-3-nitropyridine * (4.33g), following the procedure of Description 19(a). Some solid precipitated from the ethyl acetate extract, so this extract, still containing residual brine, was evaporated to dryness, washed with water, and dried. This gave the title compound (4.83g) as a brown powder.

55

NMR (δ) ($\text{DMSO}-d_6$): 2.7 (2H,t), 3.3 (2H,t), 7.7 (1H,d), 8.65 (1H,d), 9.25 (1H,s), 12.55 (1H,b).

* J. Med. Chem. 32 (11), 2477 (1989).

b) 3,4-Dihydro-4-oxo-2H-pyrido[4,3-b]-1,4-thiazepine

This material is formed from 3-((3-nitro-4-pyridyl)thio)propanoic acid, following the procedures of Descriptions 19(b) and 19(c).

c) 3,4-Dihydro-4-oxo-2H-pyrido[4,3-b]-1,4-thiazepine

This material is formed from 3,4-dihydro-4-oxo-2 H -pyrido[4,3-b]-1,4-thiazepine following the procedures of Descriptions 2(a) and 7(d).

Description 22a) 2,3-Dihydro-3-oxo-4H-pyrido[4,3-b]-1,4-oxazine

This material is formed from 4-chloro-3-nitropyridine, following the procedures of Descriptions 20(a)-(c).

b) 2,3-Dihydro-3-oxo-4H-pyrido[4,3-b]-1,4-oxazine-4-propanoic acid

This material is formed from 2,3-dihydro-3-oxo-4 H -pyrido[4,3-b]-1,4-oxazine, following the procedures of Descriptions 2(a) and 7(d).

Example 130 2,3-Dihydrofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA isobutylamide.0.5H₂O

This material was formed from 2,3-dihydrofuro[3,2-b]pyridine-3-acetic acid (Description 1) (0.074g) and Phe-Leu-ACHPA isobutylamide (Description 5(c)) (0.15g), following the procedure of Description 3(a). The crude product was purified by chromatography on silica gel using methanol/chloroform (0-4% methanol, gradient) to afford the title compound (0.13g) as a cream solid.

NMR (δ) (CD₃OD): 0.7-1.9(29H,m), 2.2-2.5(3H,m), 2.7-3.25(5H,m), 3.6-4.1(4H,m), 4.2-4.45(2H,m), 4.5-4.75-(2H,m), 7.1-7.35(7H,m), 7.9(1H,m).

Analysis: C₃₉H₅₇N₅O₆.0.5H₂O requires C, 66.8; H,8.3, N, 10.0%. Found: C,66.8; H,8.4; N,9.7%.

M.S. (m/z) (FAB) (M + 1) = 692 (consistent with m.w. = 691).

Example 245 2,3-Dihydro-4-oxofuro[3,2-]pyridine-3-acetyl-Phe-Leu-ACHPA isobutylamide

To a stirred solution of 2,3-dihydrofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA isobutylamide (Example 1) (0.064g) in dry dichloromethane (1ml) at 0 °C was added 80% 3-chloroperoxybenzoic acid (MCPBA) (0.020g) in three portions over 5 mins. After stirring at room temperature for 18h further 80% MCPBA (0.002g) was added and stirring continued for 1h. The solvent was evaporated in vacuo and the residue purified by chromatography on silica gel using methanol/chloroform (0-6% methanol, gradient) to afford the title compound (0.058g) as a white solid.

M.S. (m/z) (FAB) (M + 1) = 708 (consistent with m.w. = 707).

Example 3

2,3-Dihydro-3-oxo-4H-pyrido[3,2-b]-1,4-oxazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide
.0.5(CHCl₃)

This material was formed from 2,3-dihydro-3-oxo-4 H -pyrido[3,2-b]-1,4-oxazine-4-propanoic acid (Description 2(b)) (0.094g) and Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide (Description 3(c)) (0.16g), following the procedure of Description 3(a). The crude product was purified by chromatography on silica gel using methanol/chloroform (0-11% methanol, gradient) to afford the title compound (0.186g) as a white solid.

NMR (δ) (DMSO-d₆): 0.8-2.0(24H,m), 2.1(2H,bm), 2.3-3.1(6H,m), 3.8(2H,bm), 3.95(2H,t), 4.10(2H,bm), 4.3- (1H,m), 4.5(1H,bm), 4.7(2H,s), 4.9(1H,d), 6.9(1H,s), 7.0(1H,q), 7.1-7.5(8H,m), 7.6(1H,s), 7.8(1H,t), 8.0(1H,d), 8.1-8.3(2H,m), 8.3(0.5H,s)

Analysis: C₄₂H₅₈N₈O₇.0.5CHCl₃ requires C,59.6; H,7.0; N,13.2% Found: C,60.0; H,7.0; N,13.2%.

M.S. (m/z) (FAB) (M + 1) = 787 (consistent with m.w. = 786).

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Example 4a

2,3-Dihydro-3-oxo-4H-pyrido[4,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide
.2H₂O

This material was prepared from 2,3-dihydro-3-oxo-4 H -pyrido[4,3-b]-1,4-thiazine-4-propanoic acid (Description 6(b)) (0.11g), following the procedure of Example 3. Purification by chromatography on silica gel using methanol/chloroform (0-11% methanol, gradient) afforded the title compound (0.07g).

NMR (δ) (DMSO-d₆): 0.7-1.9 (24H,m), 2.1 (2H,m), 2.4 (2H,m), 2.7-3.1 (4H,m), 3.55 (2H,s), 3.8-4.1 (1H,m), 4.9 (1H,d), 6.9 (1H,s), 7.1-7.3 (7H,m), 7.45 (1H,d), 7.6 (1H,s), 7.75 (1H,t), 8.1-8.2 (2H,m), 8.3 (1H,d), 8.7 (1H,s).

Analysis: C₄₂H₅₈N₈O₆S.2H₂O requires C,60.1; H,7.5; N,13.4%. Found: C,60.1; H,7.1; N,13.0%.

M.S. (m/z) (FAB) (M + 1) = 803 (consistent with m.w. = 802).

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Example 4b

2,3-Dihydro-3-oxo-4H-pyrido[4,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide
.HCl

The corresponding free base (Example 4(a)) (0.03g) was dissolved in methanol (1 ml), acidified with a few drops of 5M hydrochloric acid, diluted with water (25 ml), and freeze-dried. This gave the title compound (0.03g) as a white powder.

NMR (δ) (DMSO-d₆): 0.7-1.0 (8H,m), 1.0-1.45 (7H,m), 1.5-1.8 (7H,m), 1.9 (2H,m), 2.1 (2H,m), 2.45 (2H,m), 2.7 (1H,dd), 3.0 (3H,m), 3.95 (m), 4.05 (m), 4.2 (2H,t), 4.3 (1H,q), 4.55 (1H,m), 7.1-7.3 (5H,m), 7.4 (1H,d), 7.6 (1H,d), 7.7 (1H,s), 7.8 (1H,s), 7.85 (1H,t), 8.2 (2H,m), 8.35 (1H,d), 8.6 (1H,s), 9.1 (1H,s), 14.4 (b).

M.S. (m/z) (FAB) (M + 1) = 803 (consistent with m.w. of free base = 802).

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Example 5

2,3-Dihydrofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide

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The material was prepared from 2,3-dihydrofuro[3,2-b]pyridine-3-acetic acid (Description 1(a)) (0.14g), following the procedure of Example 3. This gave the title compound (0.25g).

NMR (δ) (DMSO-d₆): 0.7-1.9 (24H,m), 2.1 (2H,m), 2.3 (1H,m), 2.55-3.1 (5H,m), 3.7-4.0 (4H,m), 4.15-4.6 (3H,m), 4.7-4.9 (1H,bm), 6.9 (1H,s), 7.0-8.3 (14H,m).

M.S. (m/z) (FAB) (M + 1) = 744 (consistent with m.w. = 743).

Example 6

2,3-Dihydro-2-oxo-1H-imidazo[4,5-b]pyridine-3-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.HCl

This material was formed from 2,3-dihydro-2-oxo-1 H -imidazo[4,5-b]pyridine-3-propanoic acid (Description 7(d)) (0.10g) and Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.2(CF₃CO₂H) (Description 9(b)) (0.40g), following the coupling procedure of Description 3(b), but substituting N,N-diisopropylethylamine (DIPEA) for triethylamine. The crude product was purified by chromatography on silica gel using methanol/chloroform (0-10% methanol, gradient), giving the free base of the title compound. This was dissolved in methanol (2 ml), a slight excess of 5M hydrochloric acid was added, and the mixture was diluted with water (30 ml) and freeze-dried, giving the title compound (0.19g).

NMR (δ) (DMSO-d₆): 0.7-1.0 (8H,m), 1.0-1.5 (7H,m), 1.5-1.85 (7H,m), 1.95 (2H,m), 2.15 (2H,m), 2.55 (2H,m), 2.8 (1H,m), 2.9-3.1 (3H,m), 3.8-4.0 (4H,m), 4.5 (1H,m), 7.0 (1H,dd), 7.1-7.4 (6H,m), 7.7 (1H,s), 7.8-8.0 (3H,m), 8.2 (1H,d), 8.3 (1H,m), 9.15 (1H,s), 11.15 (1H,s), 14.55 (1H,b).

M.S. (m/z) (FAB) (M + 1) = 772 (consistent with m.w. of free base = 771).

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Example 72,3-Dihydro-2-oxo-1H-imidazo[4,5-b]pyridine-1-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.HCl

20

This material was formed from 2,3-dihydro-2-oxo-1 H -imidazo[4,5-b]pyridine-1-propanoic acid (Description 8(d)) (0.10g), following the procedure of Example 6. This gave the title compound (0.20g) as a white powder.

NMR (δ) (DMSO-d₆): 0.7-1.8 (22H,m), 1.95 (2H,m), 2.1 (2H,m), 2.5 (m), 2.7 (1H,dd), 3.0 (3H,m), 3.85 (4H,m), 4.2 (m), 4.3 (m), 4.5 (1H,m), 6.95 (1H,dd), 7.1-7.4 (7H,m), 7.7 (1H,s), 7.8 (1H,s), 7.9 (2H,m), 8.2-8.4 (2H,m), 9.1 (1H,s), 11.5 (1H,b), 14.55 (1H,b).

M.S. (m/z) (FAB) (M + 1) = 772 (consistent with m.w. of free base = 771).

30 Example 82,3-Dihydro-4-oxofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.2HCl

This material was formed from 2,3-dihydrofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide (Example 5) (0.31g), following the procedure of Example 2. After purification by chromatography, acidification and freeze-drying was performed as described in Example 6. This gave the title compound (0.12g).

NMR (δ) (DMSO-d₆): 0.7-1.85 (22H,m), 1.9 (2H,m), 2.1 (2H,m), 2.25-2.45 (1H,m), 2.75 (1H,m), 3.05 (4H,m), 3.8-4.1 (3H,m), 4.2-4.4 (3H,m), 4.55 (2H,m), 6.9 (1H,m), 7.1-7.9 (12H,m), 8.2-8.5 (2H,m), 9.15 (1H,s), 14.7 (1H,b).

M.S. (m/z) (FAB) (M + 1) = 760 (consistent with m.w. of free base = 759).

45 Example 92,3-Dihydro-1,1,3-trioxo-4H-pyrido[4,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide .HCl

50

This material was formed from 2,3-dihydro-1,1,3-trioxo-4 H -pyrido[4,3-b]-1,4-thiazine-4-propanoic acid (Description 10(c)) (0.058g), following the procedure of Example 6. This gave the title compound (0.098g) as a fluffy white powder.

NMR (δ) (DMSO-d₆): 0.7-1.0 (8H,m), 1.0-1.5 (7H,m), 1.5-1.8 (7H,m), 1.9 (2H,m), 2.1 (2H,m), 2.45 (2H,m), 2.7 (1H,dd), 3.05 (3H,m), 3.85 (2H,bs), 4.0-4.25 (4H,m), 4.3 (1H,q), 4.6 (1H,m), 4.95 (2H,m), 7.1-7.3 (6H,m), 7.35 (1H,d), 7.7 (1H,s), 7.8 (1H,s), 7.85 (2H,m), 8.25 (1H,d), 8.4 (1H,d), 8.65 (1H,d), 8.95 (1H,s), 9.1 (1H,s), 14.45 (1H,b).

M.S. (m/z) (FAB)(M + 1) = 835 (consistent with m.w. of free base = 834).

Example 10

2,3-Dihydro-3-oxo-4H-pyrido[2,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide
 5 .HCl

This material was formed from 2,3-dihydro-3-oxo-4 H -pyrido[2,3-b]-1,4-thiazine-4-propanoic acid (Description 11(c)) (0.12g), following the procedure of Example 6. This gave the title compound (0.21g) as a white solid.

10 NMR (δ) (DMSO d_6): 0.7-1.0 (8H,m), 1.0-1.5 (7H,m), 1.5-1.8 (7H,m), 1.9 (2H,m), 2.1 (2H,m), 2.4 (2H,m), 2.7 (1H,dd), 3.05 (3H,m), 3.65 (2H,s), 3.85 (3H,m), 4.0 (1H,m), 4.2 (2H,t), 4.3 (1H,q), 4.55 (1H,m), 7.1-7.3 (6H,m), 7.45 (1H,d), 7.7 (2H,m), 7.8 (1H,s), 7.9 (1H,m), 8.15 (1H,d), 8.25 (1H,d), 8.4 (1H,d), 9.15 (1H,s), 14.65 (1H,b).
 M.S. (m/z) (FAB) (M + 1) = 803 (consistent with m.w. of free base = 802).

15

Example 11

2,3-Dihydro-1,1,3-trioxo-4H-pyrido[2,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-
 20 propylamide .HCl.1.5H $_2$ O

This material was formed from 2,3-dihydro-1,1,3-trioxo-4 H -pyrido[2,3-b]-1,4-thiazine-4-propanoic acid (Description 12(b)) (0.15g), following the procedure of Example 6. This gave the title compound (0.23g) as a white solid.

25 NMR (δ) (DMSO d_6): 0.7-1.0 (8H,m), 1.0-1.5 (7H,m), 1.5-1.8 (7H,m), 1.9 (2H,m), 2.15 (2H,m), 2.4 (2H,m), 2.75 (1H,dd), 3.05 (3H,m), 3.85 (2H,bs), 3.95 (1H,m), 4.05 (1H,m), 4.2 (2H,t), 4.3 (1H,q), 4.55 (1H,m), 4.9 (2H,m), 7.1-7.3 (6H,m), 7.4 (1H,d), 7.7 (1H,s), 7.75-7.9 (3H,m), 8.05 (1H,m), 8.25 (1H,d), 8.4 (1H,d), 8.5 (1H,d), 9.1 (1H,s), 14.5 (1H,b).

Analysis: C $_{42}$ H $_{58}$ N $_8$ O $_8$ S.HCl.1.5H $_2$ O requires C,56.1; H,7.0; N,12.5%. Found: C,56.0; H,6.6; N,12.2%.

30 M.S. (m/z) (FAB) (M + 1) = 835 (consistent with m.w. of free base = 834).

Example 12

35

6,7-Dihydro-5-oxo-5H-pyrrolo[3,4-b]pyridine-6-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.HCl

This material was formed from 6,7-dihydro-5-oxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoyl-Phe-Leu-ACHPA (Description 13(d)) (0.15g), following the procedure of Example 6. This gave the title compound (0.14g).

40 NMR (δ) (DMSO d_6): 0.6-1.0 (8H,m), 1.0-1.4 (6H,m), 1.5-1.8 (8H,m), 1.9 (2H,t), 2.1 (3H,m), 2.7-3.1 (3H,m), 3.4 (1H,m), 3.8 (2H,m), 4.1-4.5 (5H,m), 4.5-5.4 (m), 7.1-7.6 (7H,m), 7.6-8.3 (6H,m), 8.7 (1H,d), 9.1 (1H,s), 14.6 (1H,b).

M.S. (m/z) (FAB) (M + 1) = 785 (consistent with m.w. of free base = 784).

45

Example 13

6,7-Dihydro-7-oxo-5H-pyrrolo[3,4-b]pyridine-6-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.HCl
 50

This material was formed from 6,7-dihydro-7-oxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoyl-Phe-Leu-ACHPA (Description 14(c)) (0.12g), following the procedure of Example 6. This gave the title compound (0.09g).

55 NMR (δ) (DMSO d_6): 0.7-1.0 (6H,m), 1.0-1.4 (6H,m), 1.4-1.8 (8H,m), 1.9 (2H,t), 2.0-2.1 (3H,m), 2.7 (1H,t), 2.9-3.1 (3H,m), 3.4-3.5 (1H,m), 3.6 (2H,m), 4.1-4.6 (6H,m), 4.6-5.1 (m), 7.1-7.4 (6H,m), 7.6 (1H,m), 7.7 (1H,s), 7.8 (1H,s), 7.9 (1H,t), 8.05 (1H,d), 8.2 (2H,m), 8.7 (1H,d), 9.1 (1H,s), 14.6 (1H,s).

M.S. (m/z) (FAB) (M + 1) = 785 (consistent with m.w. of free base = 784).

Example 146,7-Dihydro-5H-7-oxo-pyrrolo[3,4-c]pyridine-6-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.2HCl

5 This material was formed from 6,7-dihydro-5 H -7-oxo-pyrrolo[3,4-c]pyridine-6-butanoyl acid.HCl (Description 15(c)) (0.19g), following the procedure of Example 6. This gave the title compound (0.28g).
 NMR (δ) (DMSO d_6): 0.65-1.0 (8H,m), 1.0-1.5 (8H,m), 1.5-1.8 (8H,m), 1.9 (2H,t), 2.0-2.2 (4H,m), 2.7-2.8 (1H,m), 2.9-3.1 (3H,m), 3.4 (2H,m), 3.8 (2H,s), 4.15-4.3 (3H,m), 4.5 (1H,m), 4.65 (2H,m), 4.9-5.9 (3H,m), 7.1-
 10 7.5 (6H,m), 7.6-8.3 (6H,m), 8.9 (1H,d), 9.05-9.2 (2H,m), 14.6 (1H,bs).
 M.S. (m/z) (FAB) (M + 1) = 785 (consistent with m.w. of free base = 784).

Example 157-Oxo-5,6,7,8-tetrahydro-8H-1,8-naphthyridine-8-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.HCl

15 This material was formed from 7-oxo-5,6,7,8-tetrahydro-8H-1,8-naphthyridine-8-butanoyl acid (Description 16(b)) (0.06g), following the procedure of Example 6. This gave the title compound (0.10g).
 NMR (δ) (DMSO d_6): 0.65-1.0 (8H,m), 1.0-1.75 (16H,m), 1.9 (2H,m), 2.0-2.2 (4H,m), 2.6 (2H,dd), 3.8 (2H,m), 3.9 (2H,m), 4.2 (4H,m), 4.5 (1H,m), 7.0 (1H,m), 7.2 (5H,m), 7.3-7.55 (1H,m), 7.6 (1H,d), 7.7 (1H,s), 7.8 (1H,s), 7.9 (1H,m), 8.1-8.3 (3H,m), 9.1 (1H,s), 14.5 (1H,b).
 M.S. (m/z) (FAB) (M + 1) = 799 (consistent with m.w. of free base = 798).

Example 166-Oxo-5,6,7,8-tetrahydro-5H-1,5-naphthyridine-5-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.HCl

20 This material was formed from 6-oxo-5,6,7,8-tetrahydro-5 H -1,5-naphthyridine-5-butanoyl acid (Description 17(b)) (0.12g), following the procedure of Example 6. This gave the title compound (0.23g).
 NMR (δ) (DMSO d_6): 0.7-0.95 (8H,m), 1.0-1.8 (16H,m), 1.9 (2H,m), 2.1 (4H,m), 2.8 (2H,m), 3.0 (2H,m), 3.35 (2H,m), 3.8 (2H,b), 4.2 (2H,m), 4.55 (1H,m), 7.0-7.3 (6H,m), 7.4-7.6 (1H,m), 7.65 (1H,s), 7.8 (2H,m), 7.85-8.1 (2H,m), 8.15-8.4 (3H,m), 9.15 (1H,d).
 M.S. (m/z) (FAB) (M + 1) = 799 (consistent with m.w. of free base = 798).

Example 17a7-Oxo-4,5,6,7-tetrahydro-6H-thieno[2,3-c]pyridine-6-pentanoyl-Phe-Leu ACHPA 3-(1-imidazolyl)-propylamide.H₂O

45 This material is formed from 7-oxo-4,5,6,7-tetrahydro-6H-thieno[2,3-c]pyridine-6-pentanoic acid (Description 18(b)) following the coupling procedure of Example 6. The crude product is purified by column chromatography on silica gel using methanol/chloroform (0-10% methanol/chloroform, gradient) to give the title compound.
 50 NMR (δ) (DMSO d_6): 0.7-1.9 (29H,m), 2.0-2.2 (4H,m), 2.7-3.2 (6H,m), 3.5 (2H,m), 3.8 (2H,m), 3.9 (2H,m), 4.3 (1H,m), 4.5 (1H,m), 4.8-4.9 (1H,m), 6.9 (1H,d), 7.0 (1H,dd), 7.1 (2H,m), 7.2-7.3 (5H,m), 7.5 (1H,d), 7.6 (1H,d), 7.7 (2H,m), 8.0-8.2 (2H,m).
 M.S. (m/z) (FAB) (M + 1) = 818 (consistent with m.w. = 817).
 Analysis: C₄₄H₆₃N₇O₆S.H₂O requires C,63.2; H,7.8; N,11.7%. Found: C,63.2; H,7.8; N,11.8%.

Example 17b

7-Oxo-4,5,6,7-tetrahydro-6H-thieno[2,3-c]pyridine-6-pentanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide .HCl.1.5H₂O

This material is formed from the corresponding free base (Example 17(a)) following the procedure of Example 4(b).

NMR (δ) (DMSO-d₆): 0.7-1.9 (29H,m), 2.1-2.3 (5H,m), 2.7-3.1 (5H,m), 3.3-3.4 (2H,m), 3.5 (2H,q), 3.9 (2H,m), 4.2-4.6 (3H,m), 7.0 (1H,m), 7.1 (1H,m), 7.2-7.3 (5H,m), 7.4 (1H,d), 7.6 (1H,d), 7.7 (1H,s), 7.7-7.8 (2H,m), 7.9 (1H,t), 8.0-8.3 (2H,m), 9.1 (1H,s), 14.5 (1H,bs).

Analysis: C₄₄H₆₃N₇O₆S.HCl .1.5H₂O requires C,59.9; H,7.6; N,11.1%. Found: C,59.8; H,7.6; N,11.2%.

Example 183,4-Dihydro-4-oxo-2H-pyrido[2,3-b]-1,4-thiazepine-5-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide

This material is formed from 3,4-dihydro-4-oxo-2 H -pyrido[2,3-b]-1,4-thiazepine-5-propanoic acid (Description 19(e)), following the procedure of Example 3.

Example 192,3-Dihydro-3-oxo-4H-pyrido[2,3-b]-1,4-oxazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide

This material is formed from 2,3-dihydro-3-oxo-4 H -pyrido[2,3-b]-1,4-oxazine-4-propanoic acid (Description 20(d)), following the procedure of Example 3.

Example 203,4-Dihydro-4-oxo-2H-pyrido[4,3-b]-1,4-thiazepine-5-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide

This material is formed from 3,4-dihydro-4-oxo-2 H -pyrido[4,3-b]-1,4-thiazepine-5-propanoic acid (Description 21(c)), following the procedure of Example 3.

Example 212,3-Dihydro-3-oxo-4H-pyrido[4,3-b]-1,4-oxazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide

This material is formed from 2,3-dihydro-3-oxo-4 H -pyrido[4,3-b]-1,4-oxazine-4-propanoic acid (Description 22(b)), following the procedure of Example 3.

Biological DataIn vitro human renin inhibition

Renin inhibitory activity was estimated as the percentage change in renin activity in human plasma in the presence and absence of compound. The source of plasma was blood taken from healthy volunteers. Renin activity was defined by the difference in angiotensin I levels between two halves of a sample, one incubated at 37° and the other at 4° for 2 h. Angiotensin I levels were measured using a ¹²⁵I- angiotensin I radioimmunoassay kit (NEN/Dupont, Stevenage). Results were calculated as the mean of at least two, duplicate determinations and IC₅₀ values were calculated by linear regression analysis of at least three

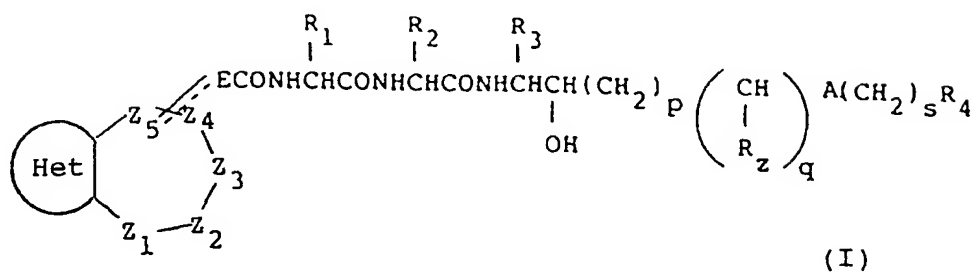
concentrations of compound.

The results were as follows:

Compound	IC ₅₀ (n- m)
E2	21
E3	26
E4	4
E10	6
E11	12
E14	14

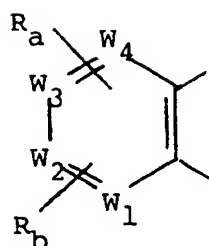
Claims

1. A compound of formula (I), or a pharmaceutically acceptable salt thereof:

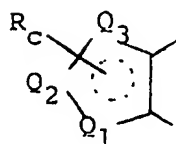


wherein

the Het ring is of sub-formula (a) or (b):



(a)



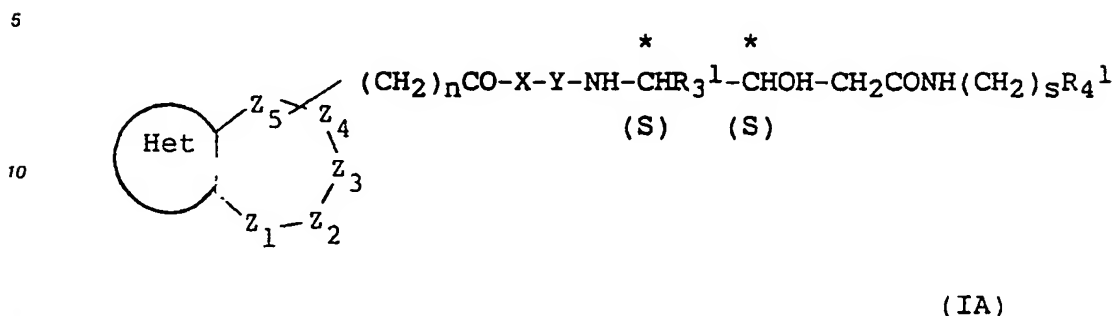
(b)

wherein

- one or two of W_1 , W_2 , W_3 and W_4 is N or NO (such that if two are NO then these are W_1 and W_4), and the others are CH, CR_a or CR_b ;
 one of Q_1 , Q_2 and Q_3 is S and the other two are CH and CR_c ;
 either Z_1 and Z_2 are absent and Z_3 , Z_4 and Z_5 and the carbon atoms to which Z_3 and Z_5 are attached, form a 5-membered non-aromatic heterocyclic ring; or
 Z_1 is absent and Z_2 , Z_3 , Z_4 , Z_5 and the carbon atoms to which Z_2 and Z_5 are attached, form a 6-membered non-aromatic heterocyclic ring; or
 Z_1 , Z_2 , Z_3 , Z_4 and Z_5 and the carbon atoms to which Z_1 and Z_5 are attached, form a 7-membered non-aromatic heterocyclic ring;
 E is absent or is $(CH_2)_n$ or $CH(CH_2)_{n-1}$ wherein n is 1 to 4;
 A is -CONH-, -NHCO-, -COO-, $-S(O)_r$ wherein r is 0, 1 or 2, or $-CH_2-$;
 p is 0, 1 or 2;
 s is 0, 1, 2, 3 or 4;
 q is 0 or 1;
 R_2 is hydrogen, C_1-6 alkyl or, when A is $-CH_2-$, hydroxy;
 R_a and R_b are independently selected from hydrogen or a substituent;
 R_1 is CH_2R_9 wherein R_9 is optionally substituted aryl or heteroaryl;
 R_2 is $CHR_{10}R_{11}$ wherein R_{10} is hydrogen or methyl and R_{11} is C_1-6 alkyl, C_3-8 cycloalkyl, optionally substituted aryl or heteroaryl, or R_{11} is amino, C_2-7 alkanoylamino, 2-oxopyrrolidinyl, 2-oxopiperidinyl or C_1-6 alkoxy carbonylamino;
 R_3 is CH_2R_{12} wherein R_{12} is C_1-6 alkyl, C_3-8 cycloalkyl or phenyl;
 R_4 is C_1-6 alkyl, C_3-8 cycloalkyl, a saturated or unsaturated heterocyclic ring linked through carbon, hydroxy, C_1-6 alkoxy, C_1-7 alkanoyloxy, amino, C_1-7 alkanoylamino, amino substituted by one or two C_1-6 alkyl groups, or C_1-6 alkylsulphonyl, carboxy, C_1-6 alkoxy carbonyl, benzyloxycarbonyl, aminocarbonyl or $CH(NHR_{13})CO_2R_{14}$ wherein R_{13} is hydrogen or C_1-6 alkanoyl and R_{14} is hydrogen or C_1-6 alkyl; or (when s is 2 to 4) R_4 is a saturated or unsaturated heterocyclic ring linked through nitrogen; and the dashed line represents an optional bond (when E is present).
 2. A compound according to claim 1 wherein Z_1 and Z_2 are absent, Z_3 is CO, Z_4 is N, Z_5 is CH and E is attached at Z_4 .
 3. A compound according to claim 1 wherein the Het ring is of sub-formula (a), W_3 is N and W_1 , W_2 and W_4 are CH, CR_a or CR_b , Z_1 is absent, Z_2 is S, Z_3 is CH_2 , Z_4 is CO, Z_5 is N and E is attached at Z_5 .
 4. A compound according to any one of claims 1 to 3 wherein one of R_a and R_b in sub-formula (b) is hydrogen and the other is CH_2NH_2 , CO_2H or $CH_2NHCH_2CO_2H$, or R_c in sub-formula (b) is hydrogen.
 5. A compound according to any one of claims 1 to 4 wherein the amino acid residue containing R_2 is Leu,

β -pyrazolylalanine or His.

6. A compound according to claim 1 of formula (IA) or a pharmaceutically acceptable salt thereof:



wherein

X is Phe;

Y is Leu or His;

R₃¹ is cyclohexylmethyl;

R₄ is C₄₋₅ alkyl and s is 0 or s is 2 to 4 and R₄¹ is carboxy or a saturated or unsaturated heterocyclic ring;

and

the remaining variables are as defined in claim 1.

7. A compound according to any one of claims 1 to 6 wherein E is (CH₂)_n and E is attached at Z₄ or Z₅.

8. A compound according to any one of claims 1 to 7 wherein the sequence from the amino acid containing R₃¹ or R₃¹ to NH(CH₂)_sR₄¹ is 4(S)-amino-5-cyclohexyl-3(S)-hydroxypentanoic acid (ACHPA).

9. A compound according to any one of claims 1 to 8 wherein s is 0 and R₄ is isobutyl or s is 2, 3 or 4 and R₄ is 1-imidazolyl, 2-imidazolyl, 4-morpholinyl, 2-oxopyrrolidin-1-yl, 2-pyridyl, 2-pyridyl-N-oxide, 3-pyridyl, hydroxyethyl, acetylamino, 4-methylpiperazin-1-yl, piperazin-1-yl or dimethylamino.

10. A compound selected from the group consisting of:

2,3-dihydrofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA isobutylamide,

2,3-dihydro-4-oxofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA isobutylamide,

2,3-dihydro-3-oxo-4 H -pyrido[3,2-b]-1,4-oxazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

2,3-dihydro-3-oxo-4 H -pyrido[4,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

2,3-dihydrofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

2,3-dihydro-2-oxo-1 H -imidazo[4,5-b]pyridine-3-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

2,3-dihydro-2-oxo-1 H -imidazo[4,5-b]pyridine-1-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

2,3-dihydro-4-oxofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

2,3-dihydro-1,1,3-trioxo-4 H -pyrido[4,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-

propylamide, 2,3-dihydro-3-oxo-4 H -pyrido[2,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-im-

imidazolyl)propylamide, 2,3-dihydro-1,1,3-trioxo-4 H -pyrido[2,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-

(1-imidazolyl)propylamide,

6,7-dihydro-5-oxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

6,7-dihydro-7-oxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

6,7-dihydro-5 H -7-oxo-pyrrolo[3,4-c]pyridine-6-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

7-oxo-5,6,7,8-tetrahydro-8 H -1,8-naphthyridine-8-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

6-oxo-5,6,7,8-tetrahydro-5 H -1,5-naphthyridine-5-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

7-oxo-4,5,6,7-tetrahydro-6 H -thieno[2,3-c]-pyridine-6-pentanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-

propylamide,

3,4-dihydro-4-oxo-2 H -pyrido[2,3-b]-1,4-thiazepine-5-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-

propylamide,

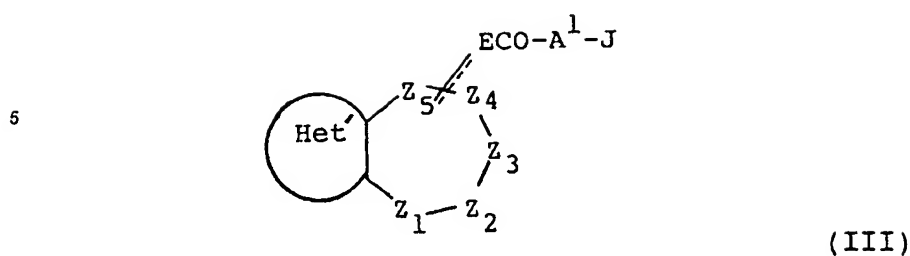
2,3-dihydro-3-oxo-4 H -pyrido[2,3-b]-1,4-oxazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,

3,4-dihydro-4-oxo-2 H -pyrido[4,3-b]-1,4-thiazepine-5-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-

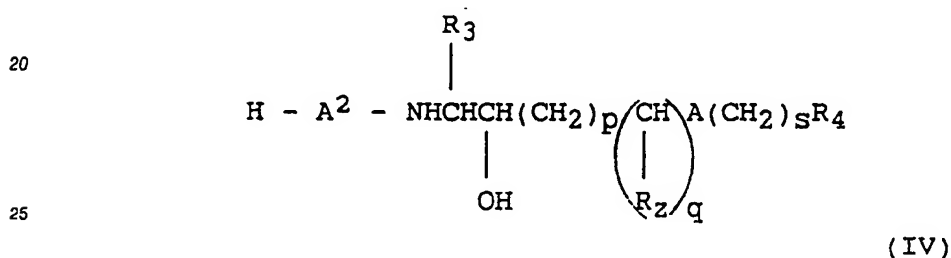
propylamide and

2,3-dihydro-3-oxo-4 H -pyrido[4,3-b]-1,4-oxazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.

11. A process for the preparation of a compound according to claim 1, which process comprises reacting a reagent of the formula (III):



15 wherein Het' is Het as defined in claim 1 but wherein R_a, R_b and R_c are R_a', R_b' and R_c', which are R_a, R_b and R_c respectively or groups or atoms convertible thereto; A¹ is absent or represents an appropriate amino acid or dipeptide unit; J is OH or a leaving group; and the remaining variables are as defined in claim 1; with a reagent of the formula (IV):



30 wherein

A² is absent or represents an appropriate amino acid or dipeptide unit, such that A¹ + A² is -NHCHR₁CONHCHR₂CO-, and the remaining variables are as defined in claim 1; and thereafter, if desired or necessary deprotecting (within A¹ or A²) of the products, and/or converting Z₁, Z₂, Z₃, Z₄ or Z₅ to other Z₁, Z₂, Z₃, Z₄ or Z₅, R_a/R_b/R_c' to R_a, R_b and/or R_c and/or forming a pharmaceutically acceptable salt thereof.

35 12. A pharmaceutical composition comprising a compound according to any one of claims 1 to 10, and a pharmaceutically acceptable carrier.

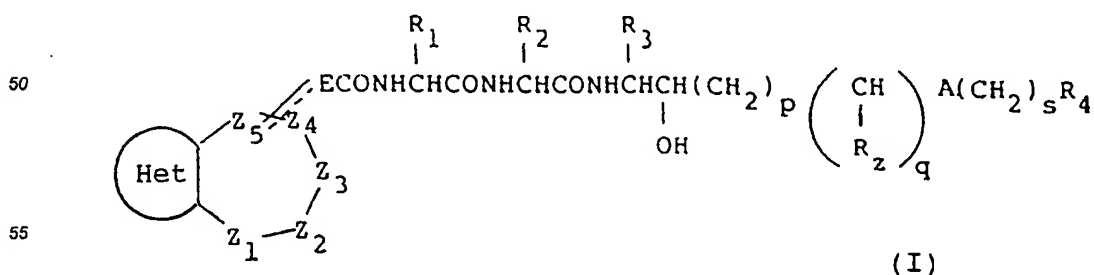
13. A compound according to any one of claims 1 to 10 for use as an active therapeutic substance.

14. A compound according to any one of claims 1 to 10 for use in the treatment of hypertension.

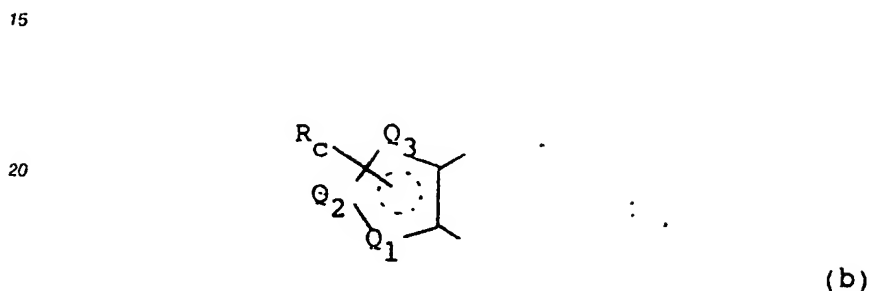
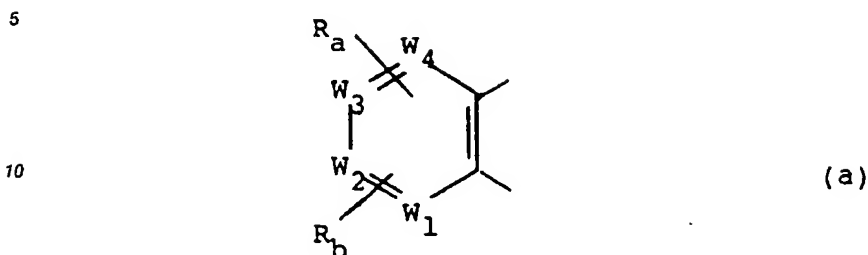
40 15. Use of a compound according to any one of claims 1 to 10 in the manufacture of a medicament for use in the treatment of hypertension.

Claims for the following Contracting State: ES.

45 1. A process for the preparation of a compound of formula (I), or a pharmaceutically acceptable salt thereof:



wherein
the Het ring is of sub-formula (a) or (b):



wherein

one or two of W_1 , W_2 , W_3 and W_4 is N or NO (such that if two are NO then these are W_1 and W_4), and the others are CH, CR_a or CR_b ;

one of Q_1 , Q_2 and Q_3 is S and the other two are CH and CR_c ;

either Z_1 and Z_2 are absent and Z_3 , Z_4 and Z_5 and the carbon atoms to which Z_3 and Z_5 are attached, form a 5-membered non-aromatic heterocyclic ring; or

Z_1 is absent and Z_2 , Z_3 , Z_4 , Z_5 and the carbon atoms to which Z_2 and Z_5 are attached, form a 6-membered non-aromatic heterocyclic ring; or

Z_1 , Z_2 , Z_3 , Z_4 and Z_5 and the carbon atoms to which Z_1 and Z_5 are attached, form a 7-membered non-aromatic heterocyclic ring;

E is absent or is $(CH_2)_n$ or $CH(CH_2)_{n-1}$ wherein n is 1 to 4;

A is $-CONH-$, $-NHCO-$, $-COO-$, $-S(O)_r-$ wherein r is 0, 1 or 2, or $-CH_2-$;

p is 0, 1 or 2;

s is 0, 1, 2, 3 or 4;

q is 0 or 1;

R_2 is hydrogen, C_{1-6} alkyl or, when A is $-CH_2-$, hydroxy;

R_a and R_b are independently selected from hydrogen or a substituent;

R_1 is CH_2R_9 wherein R_9 is optionally substituted aryl or heteroaryl;

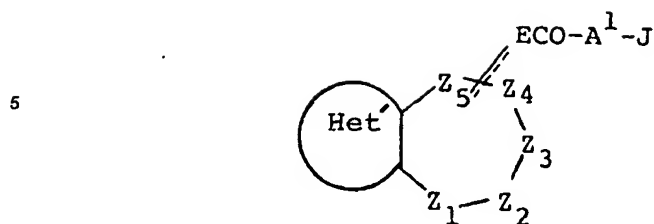
R_2 is $CHR_{10}R_{11}$ wherein R_{10} is hydrogen or methyl and R_{11} is C_{1-6} alkyl, C_{3-8} cycloalkyl, optionally substituted aryl or heteroaryl, or R_{11} is amino, C_{2-7} alkanoylamino, 2-oxopyrrolidinyl, 2-oxopiperidinyl or C_{1-5} alkoxy-carbonylamino;

R_3 is CH_2R_{12} wherein R_{12} is C_{1-6} alkyl, C_{3-8} cycloalkyl or phenyl;

R_4 is C_{1-6} alkyl, C_{3-8} cycloalkyl, a saturated or unsaturated heterocyclic ring linked through carbon, hydroxy, C_{1-6} alkoxy, C_{1-7} alkanoyloxy, amino, C_{1-7} alkanoylamino, amino substituted by one or two C_{1-6} alkyl groups, or C_{1-6} alkylsulphonyl, carboxy, C_{1-6} alkoxy-carbonyl, benzyloxy-carbonyl, aminocarbonyl or $CH(NHR_{13})CO_2R_{14}$ wherein R_{13} is hydrogen or C_{1-6} alkanoyl and R_{14} is hydrogen or C_{1-6} alkyl; or (when s is 2 to 4) R_4 is a saturated or unsaturated heterocyclic ring linked through nitrogen; and

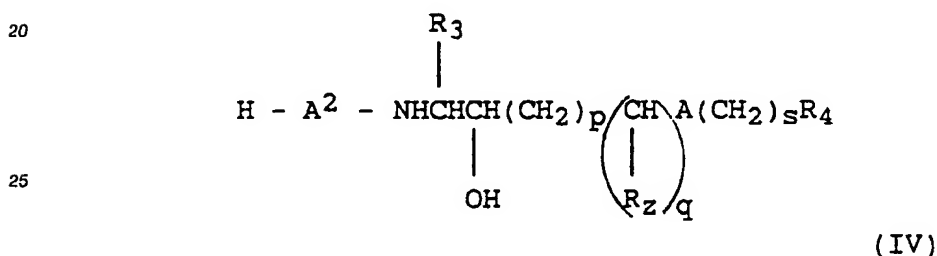
the dashed line represents an optional bond (when E is present);

which process comprises reacting a reagent of the formula (III):



(III)

15 wherein Het' is Het as defined in claim 1 but wherein R_a, R_b and R_c are R_a', R_b' and R_c', which are R_a, R_b and R_c respectively or groups or atoms convertible thereto; A' is absent or represents an appropriate amino acid or dipeptide unit; J is OH or a leaving group; and the remaining variables are as defined with a reagent of the formula (IV):



30 wherein

A² is absent or represents an appropriate amino acid or dipeptide unit, such that A¹ + A² is -NHCHR₁CONHCHR₂CO-, and the remaining variables are as defined and thereafter, if desired or necessary deprotecting (within A¹ or A²) of the products, and/or converting Z₁, Z₂, Z₃, Z₄ or Z₅ to other Z₁, Z₂, Z₃, Z₄ or Z₅, R_a/R_b/R_c to R_a, R_b and/or R_c and/or forming a pharmaceutically acceptable salt thereof.

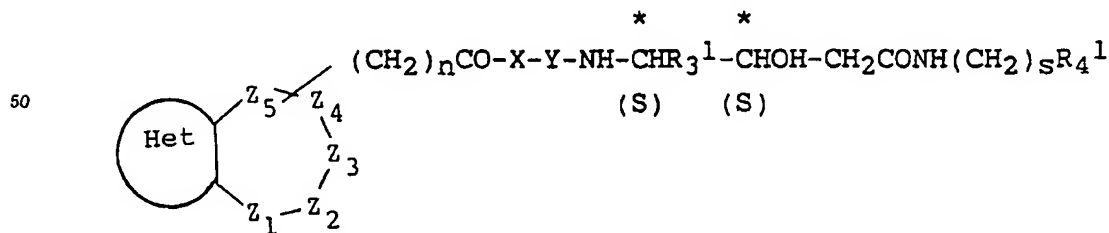
35 2. A process according to claim 1 wherein Z₁ and Z₂ are absent, Z₃ is CO, Z₄ is N, Z₅ is CH and E is attached at Z₄.

3. A process according to claim 1 wherein the Het ring is of sub-formula (a), W₃ is N and W₁, W₂ and W₄ are CH, CR_a or CR_b, Z₁ is absent, Z₂ is S, Z₃ is CH₂, Z₄ is CO, Z₅ is N and E is attached at Z₅.

40 4. A process according to any one of claims 1 to 3 wherein one of R_a and R_b in sub-formula (b) is hydrogen and the other is CH₂NH₂, CO₂H or CH₂NHCH₂CO₂H, or R_c in sub-formula (b) is hydrogen.

5. A process according to any one of claims 1 to 4 wherein the amino acid residue containing R₂ is Leu, β-pyrazolylalanine or His.

6. A process according to claim 1 for the preparation of a compound of formula (IA) or a pharmaceutically acceptable salt thereof:



(IA)

wherein

X is Phe;

Y is Leu or His;

R₃' is cyclohexylmethyl;

- 5 R₄' is C₄₋₅ alkyl and s is 0 or s is 2 to 4 and R₄' is carboxy or a saturated or unsaturated heterocyclic ring; and

the remaining variables are as defined in claim 1.

7. A process according to any one of claims 1 to 6 wherein E is (CH₂)_n and E is attached at Z₄ or Z₅.

8. A process according to any one of claims 1 to 7 wherein the sequence from the amino acid containing

- 10 R₃ or R₃' to NH(CH₂)_sR₄, is 4(S)-amino-5-cyclohexyl-3(S)-hydroxypentanoic acid (ACHPA).

9. A process according to any one of claims 1 to 8 wherein s is 0 and R₄ is isobutyl or s is 2, 3 or 4 and R₄ is 1-imidazolyl, 2-imidazolyl, 4-morpholinyl, 2-oxopyrrolidin-1-yl, 2-pyridyl, 2-pyridyl-N-oxide, 3-pyridyl, hydroxyethyl, acetyl amino, 4-methylpiperazin-1-yl, piperazin-1-yl or dimethylamino.

10. A process for the preparation of a compound selected from the group consisting of:

- 15 2,3-dihydrofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA isobutylamide,
2,3-dihydro-4-oxofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA isobutylamide,
2,3-dihydro-3-oxo-4 H -pyrido[3,2-b]-1,4-oxazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
2,3-dihydro-3-oxo-4 H -pyrido[4,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-propylamide,
2,3-dihydrofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
20 2,3-dihydro-2-oxo-1 H -imidazo[4,5-b]pyridine-3-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
2,3-dihydro-2-oxo-1 H -imidazo[4,5-b]pyridine-1-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
2,3-dihydro-4-oxofuro[3,2-b]pyridine-3-acetyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
2,3-dihydro-1,1,3-trioxo-4 H -pyrido[4,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-
propylamide,
25 2,3-dihydro-3-oxo-4 H -pyrido[2,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
2,3-dihydro-1,1,3-trioxo-4 H -pyrido[2,3-b]-1,4-thiazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-
propylamide,
6,7-dihydro-5-oxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
6,7-dihydro-7-oxo-5 H -pyrrolo[3,4-b]pyridine-6-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
30 6,7-dihydro-5 H -7-oxo-pyrrolo[3,4-c]pyridine-6-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
7-oxo-5,6,7,8-tetrahydro-8 H -1,8-naphthyridine-8-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
6-oxo-5,6,7,8-tetrahydro-5 H -1,5-naphthyridine-5-butanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
7-oxo-4,5,6,7-tetrahydro-6 H -thieno[2,3-c]-pyridine-6-pentanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-
propylamide,
35 3,4-dihydro-4-oxo-2 H -pyrido[2,3-b]-1,4-thiazepine-5-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-
propylamide,
2,3-dihydro-3-oxo-4 H -pyrido[2,3-b]-1,4-oxazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide,
3,4-dihydro-4-oxo-2 H -pyrido[4,3-b]-1,4-thiazepine-5-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)-
propylamide and
40 2,3-dihydro-3-oxo-4 H -pyrido[4,3-b]-1,4-oxazine-4-propanoyl-Phe-Leu-ACHPA 3-(1-imidazolyl)propylamide.
11. Use of a compound prepared according to a process in any one of claims 1 to 10, in the manufacture of
a medicament for use in the treatment of hypertension.

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